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OM protein - protein search, using sw model

Run on: September 18, 2004, 02:28:24 ; Search time 42.4286 Seconds
(without alignments)
39.956 Million cell updates/sec

Title: US-10-029-926B-8

Perfect score: 29

Sequence: 1 MRAPVI 6

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_29Jan04.*

1: Geneseqp1980s.*

2: Geneseqp1990s.*

3: Geneseqp2000s.*

4: Geneseqp2001s.*

5: Geneseqp2002s.*

6: Geneseqp2003as.*

7: Geneseqp2003bs.*

8: Geneseqp2004s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	29	100.0	6	2 AAW77197	AAW77197 Pharmaceu
2	29	100.0	6	5 ABG78133	ABG78133 Human Fv
3	29	100.0	6	5 ABG91824	ABG91824 Human ant
4	29	100.0	246	5 ABG78329	ABG78329 Human Fv
5	29	100.0	246	5 ABG92026	ABG92026 Antibody
6	29	100.0	256	5 ABG78334	ABG78334 Human Fv
7	29	100.0	256	5 ABG92025	ABG92025 Antibody
8	29	100.0	266	5 ABG92020	ABG92020 Human ant
9	29	100.0	277	5 ABG78150	ABG78150 Human Fv
10	29	100.0	277	5 ABG91841	ABG91841 Human ant
11	29	100.0	464	5 ABG78151	ABG78151 Human Fv
12	29	100.0	464	5 ABG92021	ABG92021 Antibody
13	29	100.0	464	5 ABG91842	ABG91842 Human ant
14	28	96.6	133	6 AAU42688	AAU42688 Propionib
15	28	96.6	133	6 ABM39207	ABM39207 Propionib
16	28	96.6	438	5 ABG14569	ABG14569 Human rho
17	28	96.6	438	7 ADA13449	ADA13449 Human rho
18	28	96.6	438	7 ADA13415	ADA13415 Human rho
19	28	96.6	470	7 ADA13459	ADA13459 Mouse rho
20	28	96.6	541	4 AAG90242	AAG90242 C glutami
21	27	93.1	116	6 AAU44808	AAU44808 Propionib
22	27	93.1	116	6 ABM41327	ABM41327 Propionib
23	26	89.7	33	4 AAM82945	AAM82945 Human imm
24	26	89.7	58	4 AAU56719	AAU56719 Propionib
25	26	89.7	58	6 ABM53238	ABM53238 Propionib

26	26	89.7	226	5 ABP73581	Abp73581 Candida a
27	26	89.7	286	4 AAG98915	Aag98915 E. coli g
28	26	89.7	286	6 ABU15292	Abu15292 Protein e
29	26	89.7	290	6 ABU28269	Abu28269 Protein e
30	26	89.7	350	2 AAY35225	Aay35225 Chlamydia
31	26	89.7	405	7 ADC87283	Adc87283 Human QPC
32	26	89.7	1449	4 ABM66060	ABM66060 Drosophil
33	25	86.2	21	2 AAY12754	Aay12754 Human S
34	25	86.2	82	5 ABP08294	Abp08294 Human ORF
35	25	86.2	86	3 AAG18780	Aag18780 Zea mays
36	25	86.2	86	4 AAM90282	Aam90282 Human imm
37	25	86.2	89	3 AAG26861	Aag26861 Zea mays
38	25	86.2	102	6 ABM65837	ABM65837 Propionib
39	25	86.2	116	5 ABP32292	ABP32292 Human ORF
40	25	86.2	120	7 ADC97061	Adc97061 E. faeciu
41	25	86.2	125	2 AAY11311	Aay11311 S. pneumo
42	25	86.2	134	4 AAU45601	AAU45601 Propionib
43	25	86.2	134	6 ABM42120	ABM42120 Propionib
44	25	86.2	158	4 AAU42587	AAU42587 Propionib
45	25	86.2	158	6 ABM39106	ABM39106 Propionib

ALIGNMENTS

RESULT 1

AAW77197

ID AAW77197 standard; peptide; 6 AA.

XX

AC AAW77197;

XX 23-NOV-1998 (first entry)

XX XX Pharmaceutically active peptide 38.

DE XX Pharmaceutically active peptide; target; organ; lymphocyte; treatment;

KW KW pharmaceutical agent; disease; radioactive isotope; imaging agent.

XX OS Synthetic.

OS OS Homo sapiens.

XX XX MO9839469-A1.

PN PN

XX XX 11-SEP-1998.

XX XX 04-MAR-1998; 98WO-US004188.

XX XX 04-MAR-1997; 97US-0039509P.

PR PR 04-MAR-1997; 97US-00810074.

XX (BIOT-) BIO-TECHNOLOGY GENERAL CORP.

XX Panet A, Hagai Y, Lazarovits J, Nimrod A, Vogel T, Levanon A;

PI Zeelon E, Belkind A, Golan I;

XX WPI; 1998-495863/42.

XX New peptide(s) binding targets in organs and lymphocytes - used for the
targetted delivery of toxins, anti-cancer drugs and cardiovascular agents
to arteries, veins, placenta, liver.

XX Claim 76; Page 99; 114pp; English.

XX Sequences shown in AAW77160 to AAW77214 and AAW79167 represent non-

CC naturally- occurring pharmacuetically active peptides. These novel
peptides specifically bind to undetermined and determined targets in
various organs and in lymphocytes. The peptides can be used in
compositions, where they can be linked to pharmaceutical agents, to treat
various diseases and conditions. The peptides or chimeric polypeptides
comprising these pharmacuetically active peptides and a second peptide
CC may be labelled with a marker (radioactive isotope, etc) to form an
imaging agent. This agent is used to bind an organ so that the organ can
be imaged

XX SQ Sequence 6 AA;

Query Match 100.0%; Score 29; DB 2; Length 6;
 Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||
 1 MRAPVI 6

Db 1 MRAPVI 6

RESULT 2
 ABG78133
 ID ABG78133 standard; peptide; 6 AA.
 XX AC ABG78133;
 XX DT 15-NOV-2002 (first entry)
 XX DE Human Fv molecule hypervariable region related peptide #8.
 XX KW Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
 KW disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
 KW lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.
 XX OS Homo sapiens.
 XX PN WO200259264-A2.
 XX PD 01-AUG-2002.
 XX PF 31-DEC-2001; 2001WO-US049440.
 XX PR 29-DEC-2000; 2000US-00751181.
 XX PA (BIOT-) BIO-TECHNOLOGY GEN CORP.
 XX PI Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
 PI Plaksin D, Peretz T;
 XX WPI; 2002-619166/66.

Novel peptide/polypeptide for cancer therapy has Fv molecule, construct or fragment, or construct of fragment with enhanced binding characteristics so as to selectively bind target cell in favor of other cells.

Claim 2; Page 76; 232pp; English.

The invention relates to a peptide or polypeptide comprising an Fv molecule, a construct or fragments or a construct of a fragment with enhanced binding characteristics which selectively and/or specifically binds to a target cell in favour of other cells, where binding is primarily determined by a first hypervariable region and Fv is a single chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in association with or attached, coupled, combined, linked or fused to a pharmaceutical agent, is useful in the manufacture of a medicament, where the medicament has activity against a diseased cell, preferably a cancer cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma, myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an acute myeloid leukaemia cell). The peptide is also useful for preparing a composition for use in inhibiting the growth of a diseased or cancer cell. This sequence represents a human Fv molecule hypervariable region related peptide of the invention

Sequence 6 AA;
 Query Match 100.0%; Score 29; DB 5; Length 6;
 Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

Db 1 MRAPVI 6

RESULT 3
 ABG91824
 ID ABG91824 standard; peptide; 6 AA.
 XX AC ABG91824;
 XX DT 04-DEC-2002 (first entry)
 XX DE Human antibody fragment #8.
 XX KW Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.
 XX OS Homo sapiens.
 XX PN WO200253700-A2.
 XX PD 11-JUL-2002.
 XX PF 31-DEC-2001; 2001WO-US049442.
 XX PR 29-DEC-2000; 2000US-00751181.
 XX PR 29-DEC-2000; 2000US-0258948P.
 XX PA (BIOT-) BIO-TECHNOLOGY GEN CORP.
 XX PI Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
 PI Szanton E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
 XX WPI; 2002-674776/72.

Novel isolated epitope present on cancer cells and important in physiological phenomena such as cell rolling, metastasis and inflammation, for treating autoimmune, inflammatory or cardiovascular diseases, and cancer.

Claim 1; Page 228; Opp; English.

The invention relates to an isolated epitope present on cancer cells and important in physiological phenomena such as cell rolling, metastasis and inflammation where the epitope is capable of being bound by an antibody, its antigen-binding fragment or its complex comprising at least one antibody or its binding fragment having a first hypervariable region. The epitopes are useful for inhibiting cell rolling, inflammation, autoimmune disease, thrombosis, restenosis, metastasis, growth and/or replication of tumour or leukaemia cells, increase in number of tumour or leukaemia cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-platelet and/or cell-platelet adhesion or aggregation, for increasing mortality of tumour or leukaemia cells, for increasing the susceptibility of diseased cells to damage by anti-disease, anti-cancer or anti-leukaemia agents, or for decreasing the number of tumour or leukaemia cells in a patient, or in the manufacture of a medicament for the above mentioned purposes. The epitopes are useful for diagnosing and treating diseases such as cancer, leukaemia, autoimmune diseases, inflammatory diseases, cardiovascular diseases such as myocardial infarction, retinopathic diseases and other diseases mediated by abnormal platelet function and diseases caused by sulphated tyrosine-dependent protein-protein interactions. This sequence represents a human antibody fragment of the invention

Sequence 6 AA;
 Query Match 100.0%; Score 29; DB 5; Length 6;
 Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

QY      1 MRAPVI 6
DB      1 MRAPVI 6

RESULT 4
ABG78329
ID      ABG78329 standard; protein; 246 AA.
AC      ABG78329;
XX
DT      15-NOV-2002 (first entry)
XX
DE      Human Fv molecule hypervariable region related peptide #204.
XX
KW      Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
KW      disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
KW      lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.
XX
OS      Homo sapiens.
XX
PN      WO200259264-A2.
XX
PD      01-AUG-2002.
XX
PF      31-DEC-2001; 2001WO-US049440.
XX
PR      29-DEC-2000; 2000US-00751181.
XX
PA      (BIOT-) BIO-TECHNOLOGY GEN CORP.
XX
PI      Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
PI      Plaksin D, Peretz T;
XX
DR      WPI; 2002-619166/66.
XX
PT      Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
PT      or fragment, or construct of fragment with enhanced binding
PT      characteristics so as to selectively bind target cell in favor of other
PT      cells.
XX
PS      Disclosure; Page 44-45; 232pp; English.
XX
CC      The invention relates to a peptide or polypeptide comprising an Fv
CC      molecule, a construct or fragments of a construct of a fragment with
CC      enhanced binding characteristics which selectively and/or specifically
CC      binds to a target cell in favour of other cells, where binding is
CC      primarily determined by a first hypervariable region and Fv is a single
CC      chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
CC      association with or attached, coupled, combined, linked or fused to a
CC      pharmaceutical agent, is useful in the manufacture of a medicament, where
CC      the medicament has activity against a diseased cell, preferably a cancer
CC      cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
CC      myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
CC      acute myeloid leukaemia cell). The peptide is also useful for preparing a
CC      composition for use in inhibiting the growth of a diseased or cancer
CC      cell. This sequence represents a human Fv molecule hypervariable region
CC      related peptide of the invention
XX
SQ      Sequence 246 AA;
      Query Match      100.0%; Score 29; DB 5; Length 246;
      Best Local Similarity 100.0%; Pred. No. 1.1e+02;
      Matches      6; Conservative      0; Mismatches      0; Indels      0; Gaps      0;

QY      1 MRAPVI 6
DB      100 MRAPVI 105

RESULT 5
ABG92026
ID      ABG92026 standard; protein; 246 AA.

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XX      ABG92026;
AC
XX      04-DEC-2002 (first entry)
DT
XX      Antibody protein #5.
DE
XX
KW      Antibody; epitope; cancer; tumour; cell rolling; inflammation;
KW      metastasis; hypervariable region; autoimmune disease; thrombosis;
KW      restenosis; leukaemia; inflammatory disease; cardiovascular disease;
KW      myocardial infarction; retinopathic disease; abnormal platelet function;
KW      sulphated tyrosine-dependent protein-protein interaction.
XX
OS      Unidentified.
XX
PN      WO200253700-A2.
XX
PD      11-JUL-2002.
XX
PF      31-DEC-2001; 2001WO-US049442.
XX
PR      29-DEC-2000; 2000US-00751181.
XX
PA      (BIOT-) BIO-TECHNOLOGY GEN CORP.
XX
PI      Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
PI      Szanton E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
XX
DR      WPI; 2002-674776/72.
XX
PT      Novel isolated epitope present on cancer cells and important in
PT      physiological phenomena such as cell rolling, metastasis and
PT      inflammation, for treating autoimmune, inflammatory or cardiovascular
PT      diseases, and cancer.
XX
PS      Disclosure; Fig 52; Opp; English.
XX
CC      The invention relates to an isolated epitope present on cancer cells and
CC      important in physiological phenomena such as cell rolling, metastasis and
CC      inflammation, where the epitope is capable of being bound by an antibody,
CC      its antigen-binding fragment or its complex comprising at least one
CC      antibody or its binding fragment having a first hypervariable region. The
CC      epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
CC      disease, thrombosis, restenosis, metastasis, growth and/or replication of
CC      tumour or leukaemia cells, increase in number of tumour or leukaemia
CC      cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
CC      platelet and/or cell-platelet adhesion or aggregation, for increasing
CC      mortality of tumour or leukaemia cells, for increasing the susceptibility
CC      of diseased cells to damage by anti-disease, anti-cancer or anti-
CC      leukaemia agents, or for decreasing the number of tumour or leukaemia
CC      cells in a patient, or in the manufacture of a medicament for the above
CC      mentioned purposes. The epitopes are useful for diagnosing and treating
CC      diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
CC      diseases, cardiovascular diseases such as myocardial infarction,
CC      retinopathic diseases and other diseases mediated by abnormal platelet
CC      function and diseases caused by sulphated tyrosine-dependent protein-
CC      protein interactions. This sequence represents an antibody protein of the
CC      invention
XX
SQ      Sequence 246 AA;
      Query Match      100.0%; Score 29; DB 5; Length 246;
      Best Local Similarity 100.0%; Pred. No. 1.1e+02;
      Matches      6; Conservative      0; Mismatches      0; Indels      0; Gaps      0;

QY      1 MRAPVI 6
DB      100 MRAPVI 105

RESULT 6
ABG78334

```


XX Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.
 XX
 OS Homo sapiens.
 XX
 FN WO200253700-A2.
 XX
 PD 11-JUL-2002.
 XX
 PF 31-DEC-2001; 2001WO-US049442.
 XX
 PR 29-DEC-2000; 2000US-00751181.
 PR 29-DEC-2000; 2000US-02589489.
 XX
 PA (BIOT-) BIO-TECHNOLOGY GEN CORP.
 XX
 PI Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
 PI Szanton E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
 XX
 DR WPI; 2002-674776/72.
 XX
 PT Novel isolated epitope present on cancer cells and important in
 PT physiological phenomena such as cell rolling, metastasis and
 PT inflammation, for treating autoimmune, inflammatory or cardiovascular
 PT diseases, and cancer.
 XX
 PS Disclosure; Page 309-310; Opp; English.
 XX
 CC The invention relates to an isolated epitope present on cancer cells and
 CC important in physiological phenomena such as cell rolling, metastasis and
 CC inflammation, where the epitope is capable of being bound by an antibody,
 CC its antigen-binding fragment or its complex comprising at least one
 CC antibody or its binding fragment having a first hypervariable region. The
 CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
 CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
 CC tumour or leukaemia cells, increase in number of tumour or leukaemia
 CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
 CC platelet and/or cell-platelet adhesion or aggregation, for increasing
 CC mortality of tumour or leukaemia cells, for increasing the susceptibility
 CC of diseased cells to damage by anti-disease, anti-cancer or anti-
 CC leukaemia agents, or for decreasing the number of tumour or leukaemia
 CC cells in a patient, or in the manufacture of a medicament for the above
 CC mentioned purposes. The epitopes are useful for diagnosing and treating
 CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
 CC diseases, cardiovascular diseases such as myocardial infarction,
 CC retinopathic diseases and other diseases mediated by abnormal platelet
 CC function and diseases caused by sulphated tyrosine-dependent protein-
 CC protein interactions. This sequence represents a human antibody fragment
 CC of the invention
 XX
 SQ Sequence 266 AA;
 XX
 Query Match 100.0%; Score 29; DB 5; Length 266;
 Best Local Similarity 100.0%; Pred. No. 1.2e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MRAPVI 6
 DB 121 MRAPVI 126
 RESULT 9
 ABG78150
 ID ABG78150 standard; protein; 277 AA.
 XX
 AC ABG78150;
 XX
 DT 15-NOV-2002 (first entry)
 XX

DE Human Fv molecule hypervariable region related peptide #25.
 XX
 KW Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
 KW disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
 KW lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.
 XX
 OS Homo sapiens.
 XX
 FN WO20025264-A2.
 XX
 PD 01-AUG-2002.
 XX
 PF 31-DEC-2001; 2001WO-US049440.
 XX
 PR 29-DEC-2000; 2000US-00751181.
 XX
 PA (BIOT-) BIO-TECHNOLOGY GEN CORP.
 XX
 PI Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
 PI Plaksin D, Peretz T;
 XX
 DR WPI; 2002-619166/66.
 XX
 PT Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
 PT or fragment, or construct of fragment with enhanced binding
 PT characteristics so as to selectively bind target cell in favor of other
 PT cells.
 XX
 PS Claim 4; Page 155-156; 232pp; English.
 XX
 CC The invention relates to a peptide or polypeptide comprising an Fv
 CC molecule, a construct or fragments or a construct of a fragment with
 CC enhanced binding characteristics which selectively and/or specifically
 CC binds to a target cell in favour of other cells, where binding is
 CC primarily determined by a first hypervariable region and Fv is a single
 CC chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
 CC association with or attached, coupled, combined, linked or fused to a
 CC pharmaceutical agent, is useful in the manufacture of a medicament, where
 CC the medicament has activity against a diseased cell, preferably a cancer
 CC cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
 CC myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
 CC acute myeloid leukaemia cell). The peptide is also useful for preparing a
 CC composition for use in inhibiting the growth of a diseased or cancer
 CC cell. This sequence represents a human Fv molecule hypervariable region
 CC related peptide of the invention
 XX
 SQ Sequence 277 AA;
 XX
 Query Match 100.0%; Score 29; DB 5; Length 277;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MRAPVI 6
 DB 121 MRAPVI 126
 RESULT 10
 ABG91841
 ID ABG91841 standard; protein; 277 AA.
 XX
 AC ABG91841;
 XX
 DT 04-DEC-2002 (first entry)
 XX
 DE Human antibody fragment #25.
 XX
 KW Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.
 XX

OS Homo sapiens.
 XX WO200253700-A2.
 PN 11-JUL-2002.
 PD
 XX 31-DEC-2001; 2001WO-US049442.
 XX
 PF 29-DEC-2000; 2000US-00751181.
 XX
 PR 29-DEC-2000; 2000US-0258948P.
 XX
 PP (BIOT-) BIO-TECHNOLOGY GEN CORP.
 XX
 PI Lazarovits J, Hagai Y, Flaksin D, Vogel T, Nimrod A, Mar-Haim H;
 PI Sztanthon E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
 XX WPI; 2002-674776/72.
 DR
 XX Novel isolated epitope present on cancer cells and important in
 PT physiological phenomena such as cell rolling, metastasis and
 PT inflammation, for treating autoimmune, inflammatory or cardiovascular
 PT diseases, and cancer.
 XX
 PS Claim 23; Page 233-234; Opp; English.
 XX
 CC The invention relates to an isolated epitope present on cancer cells and
 CC important in physiological phenomena such as cell rolling, metastasis and
 CC inflammation, where the epitope is capable of being bound by an antibody,
 CC its antigen-binding fragment or its complex comprising at least one
 CC antibody or its binding fragment having a first hypervariable region. The
 CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
 CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
 CC tumour or leukaemia cells, increase in number of tumour or leukaemia
 CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
 CC platelet and/or cell-platelet adhesion or aggregation, for increasing
 CC mortality of tumour or leukaemia cells, for increasing the susceptibility
 CC of diseased cells to damage by anti-disease, anti-cancer or anti-
 CC leukaemia agents, or for decreasing the number of tumour or leukaemia
 CC cells in a patient, or in the manufacture of a medicament for the above
 CC mentioned purposes. The epitopes are useful for diagnosing and treating
 CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
 CC diseases, cardiovascular diseases such as myocardial infarction,
 CC retinopathic diseases and other diseases mediated by abnormal platelet
 CC function and diseases caused by sulphated tyrosine-dependent protein-
 CC protein interactions. This sequence represents a human antibody fragment
 CC of the invention
 XX
 SQ Sequence 277 AA;
 Query Match 100.0%; Score 29; DB 5; Length 277;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MRAPVI 6
 Db 121 MRAPVI 126
 |||||
 RESULT 11
 ABG78151
 ID ABG78151 standard; protein; 464 AA.
 XX
 AC ABG78151;
 XX
 DT 15-NOV-2002 (first entry)
 XX
 DE Human Fv molecule hypervariable region related peptide #26.
 XX
 XX Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
 KW disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
 KW lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.
 XX
 OS Homo sapiens.

XX WO200259264-A2.
 PN
 XX
 PD 01-AUG-2002.
 XX
 PF 31-DEC-2001; 2001WO-US049440.
 XX
 PR 29-DEC-2000; 2000US-00751181.
 XX
 PP (BIOT-) BIO-TECHNOLOGY GEN CORP.
 XX
 PI Hagai Y, Lazarovits J, Guy R, Lipschitz O, Sztanton E, Levanon A;
 PI Flaksin D, Peretz T;
 XX WPI; 2002-619166/66.
 DR N-PSDB; ABS63405.
 XX
 XX Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
 PT or fragment, or construct of fragment with enhanced binding
 PT characteristics so as to selectively bind target cell in favor of other
 PT cells.
 XX
 PS Claim 109; Page 93-94; 232pp; English.
 XX
 CC The invention relates to a peptide or polypeptide comprising an Fv
 CC molecule, a construct or fragments or a construct of a fragment with
 CC enhanced binding characteristics which selectively and/or specifically
 CC binds to a target cell in favour of other cells, where binding is
 CC primarily determined by a first hypervariable region and Fv is a single
 CC chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
 CC association with or attached coupled, combined, linked or fused to a
 CC pharmaceutical agent, is useful in the manufacture of a medicament, where
 CC the medicament has activity against a diseased cell, preferably a cancer
 CC cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
 CC myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
 CC acute myeloid leukaemia cell). The peptide is also useful for preparing a
 CC composition for use in inhibiting the growth of a diseased or cancer
 CC cell. This sequence represents a human Fv molecule hypervariable region
 CC related peptide of the invention
 XX
 SQ Sequence 464 AA;
 Query Match 100.0%; Score 29; DB 5; Length 464;
 Best Local Similarity 100.0%; Pred. No. 2.2e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MRAPVI 6
 Db 118 MRAPVI 123
 |||||
 RESULT 12
 ABG92021
 ID ABG92021 standard; protein; 464 AA.
 XX
 AC ABG92021;
 XX
 DT 04-DEC-2002 (first entry)
 XX
 DE Antibody protein #1.
 XX
 KW Antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.
 XX
 OS Unidentified.
 XX
 PN WO200253700-A2.
 XX
 PD 11-JUL-2002.
 XX

```

PF 31-DEC-2001; 2001WO-US049442.
XX
XX 29-DEC-2000; 2000US-00751181.
PR 29-DEC-2000; 2000US-0258948P.
XX
XX (BIOT-) BIO-TECHNOLOGY GEN CORP.
PA
XX Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
PI Szanthon E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
XX
XX WPI; 2002-674776/72.
DR N-PSDB; ABS67759.
XX
XX Novel isolated epitope present on cancer cells and important in
PI physiological phenomena such as cell rolling, metastasis and
PT inflammation, for treating autoimmune, inflammatory or cardiovascular
PT diseases, and cancer.
XX
XX Example 5; Fig 48A; Opp; English.
XX
XX The invention relates to an isolated epitope present on cancer cells and
CC important in physiological phenomena such as cell rolling, metastasis and
CC inflammation, where the epitope is capable of being bound by an antibody,
CC its antigen-binding fragment or its complex comprising at least one
CC antibody or its binding fragment having a first hypervariable region. The
CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
CC tumour or leukaemia cells, increase in number of tumour or leukaemia
CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
CC platelet and/or cell-platelet adhesion or aggregation, for increasing
CC mortality of tumour or leukaemia cells, for increasing the susceptibility
CC of diseased cells to damage by anti-disease, anti-cancer or anti-
CC leukaemia agents, or for decreasing the number of tumour or leukaemia
CC cells in a patient, or in the manufacture of a medicament for the above
CC mentioned purposes. The epitopes are useful for diagnosing and treating
CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
CC diseases, cardiovascular diseases such as myocardial infarction,
CC retinopathic diseases and other diseases mediated by abnormal platelet
CC function and diseases caused by sulphated tyrosine-dependent protein-
CC protein interactions. This sequence represents an antibody protein of the
CC invention
XX
SQ Sequence 464 AA;
Query Match 100.0%; Score 29; DB 5; Length 464;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MRAPVI 6
Db 118 MRAPVI 123
RESULT 13
ABG91842
ID ABG91842 standard; protein; 464 AA.
XX
XX ABG91842;
AC
XX
XX 04-DEC-2002 (first entry)
DT
XX
XX Human antibody fragment #26.
DE
XX
XX Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
KW metastasis; hypervariable region; autoimmune disease; thrombosis;
KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
KW myocardial infarction; retinopathic disease; abnormal platelet function;
KW sulphated tyrosine-dependent protein-protein interaction.
XX
XX Homo sapiens.
OS
XX
XX WO200253700-A2.
PN
XX

```

```

PD 11-JUL-2002.
XX
XX 31-DEC-2001; 2001WO-US049442.
XX
XX 29-DEC-2000; 2000US-00751181.
PR 29-DEC-2000; 2000US-0258948P.
XX
XX (BIOT-) BIO-TECHNOLOGY GEN CORP.
PA
XX Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
PI Szanthon E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
XX
XX WPI; 2002-674776/72.
DR
XX
XX Novel isolated epitope present on cancer cells and important in
PI physiological phenomena such as cell rolling, metastasis and
PT inflammation, for treating autoimmune, inflammatory or cardiovascular
PT diseases, and cancer.
XX
XX Disclosure; Page 234-235; Opp; English.
XX
XX The invention relates to an isolated epitope present on cancer cells and
CC important in physiological phenomena such as cell rolling, metastasis and
CC inflammation, where the epitope is capable of being bound by an antibody,
CC its antigen-binding fragment or its complex comprising at least one
CC antibody or its binding fragment having a first hypervariable region. The
CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
CC tumour or leukaemia cells, increase in number of tumour or leukaemia
CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
CC platelet and/or cell-platelet adhesion or aggregation, for increasing
CC mortality of tumour or leukaemia cells, for increasing the susceptibility
CC of diseased cells to damage by anti-disease, anti-cancer or anti-
CC leukaemia agents, or for decreasing the number of tumour or leukaemia
CC cells in a patient, or in the manufacture of a medicament for the above
CC mentioned purposes. The epitopes are useful for diagnosing and treating
CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
CC diseases, cardiovascular diseases such as myocardial infarction,
CC retinopathic diseases and other diseases mediated by abnormal platelet
CC function and diseases caused by sulphated tyrosine-dependent protein-
CC protein interactions. This sequence represents a human antibody fragment
CC of the invention
XX
SQ Sequence 464 AA;
Query Match 100.0%; Score 29; DB 5; Length 464;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MRAPVI 6
Db 118 MRAPVI 123
RESULT 14
AAU42688
ID AAU42688 standard; protein; 133 AA.
XX
XX AAU42688;
AC
XX
XX 27-FEB-2002 (first entry)
DT
XX
XX Propionibacterium acnes immunogenic protein #3594.
DE
XX
XX SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
KW dermatological; osteopathic; neuroprotectant.
XX
XX Propionibacterium acnes.
OS
XX
XX WO200181581-A2.
PN
XX

```

PD 01-NOV-2001.
 XX 20-APR-2001; 2001WO-US012865.
 PF XX
 XX 21-APR-2000; 2000US-0199047P.
 PR 02-JUN-2000; 2000US-0208641P.
 PR 07-JUL-2000; 2000US-0216747P.
 XX (CORI-) CORIXA CORP.
 XX Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;
 PI L'maisonneuve J, Zhang Y, Jen S, Carter D;
 PI WPI; 2001-616774/71.
 DR N-PSDB; AAS59518.
 XX Propionibacterium acnes polypeptides and nucleic acids useful for
 PT vaccinating against and diagnosing infections, especially useful for
 PT treating acne vulgaris.
 XX Example 1; SEQ ID NO 3883; 1069pp; English.
 PS Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic
 CC polypeptides. The proteins and their associated DNA sequences are used in
 CC the treatment, prevention and diagnosis of medical conditions caused by
 CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,
 CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.
 CC P. acnes is also involved in infections of bone, joints and the central
 CC nervous system, however it is particularly involved in the inflammatory
 CC lesions associated with acne vulgaris. A method for detecting the
 CC presence or absence of P. acnes in a patient comprises contacting a
 CC sample with a binding agent that binds to the proteins of the invention
 CC and determining the amount of bound protein in the sample. The
 CC polypeptides may be used as antigens in the production of antibodies
 CC specific for P. acnes proteins. These antibodies can be used to
 CC downregulate expression and activity of P. acnes polypeptides and
 CC therefore treat P. acnes infections. The antibodies may also be used as
 CC diagnostic agents for determining P. acnes presence, for example, by
 CC enzyme linked immunosorbent assay (ELISA). Note: The sequence data for
 CC this patent did not form part of the printed specification, but was
 CC obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 133 AA;
 Query Match 96.6%; Score 28; DB 4; Length 133;
 Best Local Similarity 83.3%; Pred. No. 1e+02;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MRAPVI 6
 Db 59 MRAPII 64
 RESULT 15
 ID ABM39207
 XX
 XX ABM39207 standard; protein; 133 AA.
 AC
 XX ABM39207;
 DT 20-OCT-2003 (first entry)
 XX
 DE Propionibacterium acnes predicted ORF-encoded polypeptide #3883.
 XX
 KW Acne vulgaris; antiseborrheic; dermatological; antibacterial;
 KW immunostimulant; immune response; vaccine.
 XX
 OS Propionibacterium acnes.
 XX
 PN WO2003033515-A1.
 XX
 PD 24-APR-2003.
 XX

PF 11-OCT-2002; 2002WO-US032727.
 XX 15-OCT-2001; 2001US-00978825.
 XX (CORI-) CORIXA CORP.
 XX Mitcham JL, Skeiky YAW, Persing DH, Bhatia A, Maisonneuve JL;
 PI Zhang Y, Wang S, Jen S, Lodes MJ, Benson DR, Jones R, Carter D;
 PI Barth B, Valliave-Douglas J;
 XX WPI; 2003-381789/36.
 DR N-PSDB; ACF64447.
 XX New Propionibacterium acnes polypeptides and polynucleotides encoding the
 PT polypeptide, useful for diagnosing, preventing or treating acne vulgaris,
 PT or for stimulating an immune response specific for a P. acnes protein.
 XX Example 1; SEQ ID NO 3883; 1481pp; English.
 PS The invention relates to an isolated polynucleotide (ACF64435-ACF64733)
 CC encoding a Propionibacterium acnes protein. The invention also relates to
 CC polypeptides encoded by the polynucleotides (ABM35624-ABM64536) and to
 CC immunogenic fragments of P. acnes polypeptides. The invention
 CC additionally encompasses expression vectors and host cells comprising a
 CC polynucleotide of the invention; antibodies against polypeptides of the
 CC invention; fusion proteins comprising a polypeptide of the invention; a
 CC method for stimulating an immune response specific for a P. acnes
 CC polypeptide and an isolated T cell population comprising P. acnes polypeptides,
 CC via this method; a vaccine composition (comprising P. acnes polypeptides,
 CC polynucleotides, antibodies, fusion proteins, T cell populations, or
 CC antigen-presenting cells that express the polypeptide); a method and kit
 CC for detecting or determining the presence or absence of P. acnes in a
 CC patient; and a method for inhibiting the development of P. acnes in a
 CC patient. The P. acnes polypeptides, polynucleotides, antibodies, fusion
 CC proteins, T cell populations or antigen-presenting cells that express the
 CC polypeptides are useful for diagnosing, preventing or treating acne
 CC vulgaris, or for stimulating an immune response specific for a P. acnes
 CC protein. The polynucleotides can also be used as probes or primers for
 CC nucleic acid hybridisation. The vaccine composition is useful for the
 CC stimulation of an immune response against P. acnes, or for treating acne,
 CC and the kit is useful for performing a diagnostic assay. The present
 CC sequence represents a polypeptide predicted to be encoded by an ORF (open
 CC reading frame) contained within the P. acnes polynucleotides of the
 CC invention. Note: The sequence data for this patent did not form part of
 CC the printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 133 AA;
 Query Match 96.6%; Score 28; DB 6; Length 133;
 Best Local Similarity 83.3%; Pred. No. 1e+02;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MRAPVI 6
 Db 59 MRAPII 64

Search completed: September 18, 2004, 03:42:51
 Job time : 46.4286 secs

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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:39:30 ; Search time 13.9286 Seconds
(without alignments)
22.239 Million cell updates/sec

Title: US-10-029-926B-8
Perfect score: 29
Sequence: 1 MRAPVI 6

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA:*
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2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
3: /cgn2_6/ptodata/2/iaa/6A_COMB.pep.*
4: /cgn2_6/ptodata/2/iaa/6B_COMB.pep.*
5: /cgn2_6/ptodata/2/iaa/PCFUS_COMB.pep.*
6: /cgn2_6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	96.6	973	4	US-09-252-991A-23944
2	26	89.7	286	4	US-09-711-164-385
3	26	89.7	350	4	US-09-198-452A-643
4	26	89.7	477	4	US-09-489-039A-8906
5	25	86.2	119	4	US-09-252-991A-28425
6	25	86.2	120	4	US-09-107-532A-6688
7	25	86.2	222	4	US-09-489-039A-8244
8	25	86.2	223	4	US-09-134-001C-6010
9	25	86.2	233	4	US-09-489-039A-7366
10	25	86.2	239	4	US-09-328-352-5866
11	25	86.2	359	4	US-09-252-991A-19964
12	25	86.2	459	4	US-09-252-991A-26783
13	25	86.2	481	4	US-09-489-039A-10515
14	25	86.2	657	4	US-09-252-991A-27682
15	25	86.2	962	4	US-09-328-352-7942
16	24	82.8	73	4	US-09-621-976-6392
17	24	82.8	76	4	US-09-636-215-575
18	24	82.8	76	4	US-09-685-166A-575
19	24	82.8	76	4	US-09-685-166A-888
20	24	82.8	82	4	US-09-134-001C-4676
21	24	82.8	89	4	US-09-134-001C-4563
22	24	82.8	93	4	US-09-938-806A-8
23	24	82.8	125	4	US-09-199-637A-403
24	24	82.8	125	4	US-09-269-410-11
25	24	82.8	162	4	US-08-801-742-3
26	24	82.8	192	1	US-08-208-008C-9
27	24	82.8	220	4	US-09-489-039A-8353

28	24	82.8	247	2	US-07-885-039B-2	Sequence 2, Appli
29	24	82.8	247	2	US-07-885-039B-8	Sequence 8, Appli
30	24	82.8	252	2	US-07-885-039B-7	Sequence 7, Appli
31	24	82.8	252	4	US-09-976-594-582	Sequence 582, App
32	24	82.8	253	4	US-09-134-001C-5435	Sequence 5435, Ap
33	24	82.8	269	4	US-09-252-991A-31653	Sequence 31653, A
34	24	82.8	324	4	US-09-489-039A-11252	Sequence 11252, A
35	24	82.8	356	4	US-09-134-001C-3408	Sequence 3408, Ap
36	24	82.8	363	3	US-08-912-560-2	Sequence 2, Appli
37	24	82.8	409	1	US-08-403-545-2	Sequence 2, Appli
38	24	82.8	409	3	US-08-404-381-2	Sequence 2, Appli
39	24	82.8	410	3	US-09-258-754-451	Sequence 451, App
40	24	82.8	411	3	US-09-258-754-448	Sequence 448, App
41	24	82.8	429	4	US-09-252-991A-30995	Sequence 30995, A
42	24	82.8	453	4	US-09-543-681A-5855	Sequence 5855, A
43	24	82.8	461	4	US-09-489-039A-8460	Sequence 8460, Ap
44	24	82.8	464	4	US-09-252-991A-16660	Sequence 16660, A
45	24	82.8	478	4	US-09-273-871A-3	Sequence 3, Appli

ALIGNMENTS

RESULT 1
US-09-252-991A-23944
; Sequence 23944, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 23944
; LENGTH: 973
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-23944

Query Match 96.6%; Score 28; DB 4; Length 973;
Best Local Similarity 83.3%; Pred. No. 2.9e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

CY 1 MRAPVI 6
Db 387 MRAPV 392

RESULT 2
US-09-711-164-385
; Sequence 385, Application US/09711164
; Patent No. 6589738
; GENERAL INFORMATION:
; APPLICANT: Forsyth, R. Allyn
; APPLICANT: Ohlsen, Kari
; APPLICANT: Zyskind, Judith
; TITLE OF INVENTION: GENES ESSENTIAL FOR MICROBIAL PROLIFERATION AND ANTISENSE THERE
; FILE REFERENCE: ELTRA.008A
; CURRENT APPLICATION NUMBER: US/09/711,164
; CURRENT FILING DATE: 2000-11-09
; PRIOR APPLICATION NUMBER: US 60/164415
; PRIOR FILING DATE: 1999-11-9
; NUMBER OF SEQ ID NOS: 469
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 385
; LENGTH: 286
; TYPE: PRT

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; ORGANISM: Escherichia coli
US-09-711-164-385

Query Match
Best Local Similarity 89.7%; Score 26; DB 4; Length 286;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 41 MRSPVI 46

RESULT 3
US-09-198-452A-643
; Sequence 643, Application US/09198452A
; Patent No. 6559294
; GENERAL INFORMATION:
; APPLICANT: Griffais, R.
; TITLE OF INVENTION: Chlamydia pneumoniae genomic sequence and polypeptides, fragments
; TITLE OF INVENTION: thereof and uses thereof, in particular for the diagnosis, prevention
; TITLE OF INVENTION: and treatment of infection
; FILE REFERENCE: 9710-003-999
; CURRENT APPLICATION NUMBER: US/09/198.452A
; CURRENT FILING DATE: 1998-11-24
; NUMBER OF SEQ ID NOS: 6849
; SEQ ID NO 643
; LENGTH: 350
; TYPE: PRT
; ORGANISM: Chlamydia pneumoniae
US-09-198-452A-643

Query Match
Best Local Similarity 89.7%; Score 26; DB 4; Length 350;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 109 LRAPVI 114

RESULT 4
US-09-489-039A-8906
; Sequence 8906, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; TITLE OF INVENTION: PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489.039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 8906
; LENGTH: 477
; TYPE: PRT
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-8906

Query Match
Best Local Similarity 89.7%; Score 26; DB 4; Length 477;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 269 MKAPVI 274

RESULT 5
US-09-252-991A-28425
; Sequence 28425, Application US/09252991A
; Patent No. 6551795

; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-28425

Query Match
Best Local Similarity 86.2%; Score 25; DB 4; Length 119;
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 63 MRAPTI 68

RESULT 6
US-09-107-532A-6688
; Sequence 6688, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; TITLE OF INVENTION: ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107.532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 6688:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 120 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHEICAL: YES
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
```

NAME/KEY: misc.feature
LOCATION: (B) LOCATION 1...120
SEQUENCE DESCRIPTION: SEQ ID NO: 6688;
US-09-107-532A-6688

Query Match 86.2%; Score 25; DB 4; Length 120;
Best Local Similarity 86.7%; Pred. No. 1.5e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPV 6
|:|:|
Db 17 MKAPV 22

RESULT 7
US-09-489-039A-8244

; Sequence 8244, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:

; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A

; CURRENT FILING DATE: 2000-01-27
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342

; SEQ ID NO 8244
; LENGTH: 222

; TYPE: PRT

; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-8244

Query Match 86.2%; Score 25; DB 4; Length 222;
Best Local Similarity 100.0%; Pred. No. 2.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPV 5
|:|:|
Db 24 MRAPV 28

RESULT 8

US-09-134-000C-6010
; Sequence 6010, Application US/09134000C

; Patent No. 6617156

; GENERAL INFORMATION:

; APPLICANT: Lynn Doucette-Stamm et al

; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; FILE REFERENCE: 032796-032
; CURRENT APPLICATION NUMBER: US/09/134,000C

; CURRENT FILING DATE: 1998-08-13

; PRIOR FILING DATE: 1997-08-15

; NUMBER OF SEQ ID NOS: 6812

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 6010

; LENGTH: 223

; TYPE: PRT

; ORGANISM: Enterococcus faecalis

US-09-134-000C-6010

Query Match 86.2%; Score 25; DB 4; Length 223;
Best Local Similarity 100.0%; Pred. No. 2.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPV 5
|:|:|
Db 127 MRAPV 131

RESULT 9

US-09-489-039A-7366
; Sequence 7366, Application US/09489039A

; Patent No. 6610836

; GENERAL INFORMATION:

; APPLICANT: Gary Breton et. al

; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001

; CURRENT APPLICATION NUMBER: US/09/489,039A

; CURRENT FILING DATE: 2000-01-27

; PRIOR FILING DATE: 1999-01-29

; NUMBER OF SEQ ID NOS: 14342

; SEQ ID NO 7366

; LENGTH: 233

; TYPE: PRT

; ORGANISM: Klebsiella pneumoniae

US-09-489-039A-7366

Query Match 86.2%; Score 25; DB 4; Length 233;
Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPV 5
|:|:|
Db 173 MRAPV 177

RESULT 10

US-09-328-352-5866
; Sequence 5866, Application US/09328352

; Patent No. 6562958

; GENERAL INFORMATION:

; APPLICANT: Gary L. Breton et al.

; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER

; FILE REFERENCE: GTC99-03PA

; CURRENT APPLICATION NUMBER: US/09/328,352

; CURRENT FILING DATE: 1999-06-04

; NUMBER OF SEQ ID NOS: 8252

; SEQ ID NO 5866

; LENGTH: 239

; TYPE: PRT

; ORGANISM: Acinetobacter baumannii

US-09-328-352-5866

Query Match 86.2%; Score 25; DB 4; Length 239;
Best Local Similarity 66.7%; Pred. No. 3.1e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPV 6
|:|:|
Db 140 MKAPV 145

RESULT 11

US-09-252-991A-19964
; Sequence 19964, Application US/09252991A

; Patent No. 6551795

; GENERAL INFORMATION:

; APPLICANT: Marc J. Ruberfield et al.

; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS

; FILE REFERENCE: 107196.136

; CURRENT APPLICATION NUMBER: US/09/252,991A

; CURRENT FILING DATE: 1999-02-18

; PRIOR APPLICATION NUMBER: US 60/074,788

; PRIOR FILING DATE: 1998-02-18

; PRIOR APPLICATION NUMBER: US 60/094,190

; PRIOR FILING DATE: 1998-07-27

; NUMBER OF SEQ ID NOS: 33142

; SEQ ID NO 19964

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; LENGTH: 359
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-19964

Query Match      86.2%; Score 25; DB 4; Length 359;
Best Local Similarity 66.7%; Pred. No. 4.8e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
      :||||:
Db      212 LRAPV 217

RESULT 12
US-09-252-991A-26783
; Sequence 26783, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 26783
; LENGTH: 459
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-26783

Query Match      86.2%; Score 25; DB 4; Length 459;
Best Local Similarity 100.0%; Pred. No. 6.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPV 5
      :||||:
Db      38 MRAPV 42

RESULT 13
US-09-489-039A-10515
; Sequence 10515, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; PRIOR FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 10515
; LENGTH: 481
; TYPE: PRT
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-10515

Query Match      86.2%; Score 25; DB 4; Length 481;
Best Local Similarity 100.0%; Pred. No. 6.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPV 5
      :||||:
Db      131 MRAPV 135

RESULT 14
US-09-252-991A-27682
; Sequence 27682, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 27682
; LENGTH: 657
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-27682

Query Match      86.2%; Score 25; DB 4; Length 657;
Best Local Similarity 66.7%; Pred. No. 9.1e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
      :||||:
Db      332 LRAPV 337

RESULT 15
US-09-328-352-7942
; Sequence 7942, Application US/09328352
; Patent No. 6562956
; GENERAL INFORMATION:
; APPLICANT: Gary L. Breton et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER
; FILE REFERENCE: GTC99-03PA
; CURRENT APPLICATION NUMBER: US/09/328,352
; CURRENT FILING DATE: 1999-06-04
; NUMBER OF SEQ ID NOS: 8252
; SEQ ID NO 7942
; LENGTH: 962
; TYPE: PRT
; ORGANISM: Acinetobacter baumannii
US-09-328-352-7942

Query Match      86.2%; Score 25; DB 4; Length 962;
Best Local Similarity 83.3%; Pred. No. 1.4e+03;
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 MRAPVI 6
      :||||:
Db      471 MRAPV 476

Search completed: September 18, 2004, 03:48:46
Job time : 15.9286 secs
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:46:39 ; Search time 238.286 Seconds
(without alignments)
8.086 Million cell updates/sec

Title: US-10-029-926B-8

Perfect score: 29

Sequence: 1 MRAPVI 6

Scoring table: BLOSUM62
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Searched: 1342398 seqs, 32113274 residues

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA.*

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9: /cgn2_6/ptodata/2/pubpaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/2/pubpaa/US09B_PUBCOMB.pep.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	29	100.0	6	12	US-10-029-926B-8
2	29	100.0	6	15	US-10-032-037B-8
3	29	100.0	6	15	US-10-029-988B-8
4	29	100.0	6	15	US-10-032-423A-8
5	29	100.0	266	15	US-10-032-037B-204
6	29	100.0	266	15	US-10-029-988B-204
7	29	100.0	266	15	US-10-032-423A-204
8	29	100.0	277	12	US-10-029-926B-25
9	29	100.0	277	15	US-10-032-037B-25
10	29	100.0	277	15	US-10-029-988B-25
11	29	100.0	277	15	US-10-032-423A-25
12	29	100.0	442	15	US-10-369-493-7871
13	29	100.0	464	12	US-10-029-926B-26
14	29	100.0	464	15	US-10-032-037B-26
15	29	100.0	464	15	US-10-029-988B-26

16 100.0 464 15 US-10-032-423A-26
17 96.6 86 12 US-10-424-599-253048
18 96.6 123 12 US-10-424-599-150638
19 96.6 438 9 US-09-908-419-2
20 96.6 438 10 US-09-907-187-2
21 96.6 438 14 US-10-056-790-2
22 96.6 438 14 US-10-056-790-36
23 96.6 470 14 US-10-056-790-46
24 96.6 541 9 US-09-738-626-3996
25 93.1 99 12 US-10-424-599-272558
26 89.7 112 16 US-10-767-701-42410
27 89.7 226 14 US-10-032-585-7418
28 89.7 286 12 US-10-282-122A-43216
29 89.7 286 14 US-10-287-274-385
30 89.7 286 15 US-10-369-493-23563
31 89.7 289 12 US-10-425-114-69148
32 89.7 290 12 US-10-282-122A-56193
33 89.7 319 12 US-10-425-114-55531
34 89.7 345 16 US-10-767-701-45307
35 89.7 350 15 US-10-289-762-643
36 89.7 405 14 US-10-017-161-2090
37 89.7 405 15 US-10-292-798-1736
38 89.7 514 12 US-10-425-114-53835
39 89.7 530 12 US-10-425-114-60637
40 89.7 1428 16 US-10-437-963-144989
41 86.2 58 16 US-10-437-963-137085
42 86.2 63 16 US-10-767-701-48548
43 86.2 75 12 US-10-424-599-196866
44 86.2 104 16 US-10-437-963-164399
45 86.2 110 16 US-10-437-963-158848

ALIGNMENTS

RESULT 1

US-10-029-926B-8
; Sequence 8, Application US/10029926B
; Publication No. US20040073011A1
; GENERAL INFORMATION:
; APPLICANT: HAGAY, et al.
; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
; FILE REFERENCE: 10793/50
; CURRENT APPLICATION NUMBER: US/10/029,926B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 203
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 6
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-926B-8

Query Match 100.0%; Score 29; DB 12; Length 6;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 MRAPVI 6
Db 1 MRAPVI 6

RESULT 2

US-10-032-037B-8
; Sequence 8, Application US/10032037B
; Publication No. US20040001822A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/44

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; CURRENT APPLICATION NUMBER: US/10/032,037B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 6
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-8

Query Match      100.0%; Score 29; DB 15; Length 6;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MEAPVI 6
   |||||
Db 1 MEAPVI 6

RESULT 3
US-10-029-988B-8
; Sequence 8, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; PRIOR FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 6
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-8

Query Match      100.0%; Score 29; DB 15; Length 6;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MEAPVI 6
   |||||
Db 1 MEAPVI 6

RESULT 4
US-10-032-423A-8
; Sequence 8, Application US/10032423A
; Publication No. US20040002450A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/45
; CURRENT APPLICATION NUMBER: US/10/032,423A
; PRIOR FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 6
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-423A-8

Query Match      100.0%; Score 29; DB 15; Length 6;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MEAPVI 6
   |||||
Db 1 MEAPVI 6

US-10-029-926b-8.rapb

; CURRENT APPLICATION NUMBER: US/10/032,037B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 6
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-204

Query Match      100.0%; Score 29; DB 15; Length 6;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MEAPVI 6
   |||||
Db 1 MEAPVI 6

RESULT 5
US-10-032-037B-204
; Sequence 204, Application US/10032037B
; Publication No. US20040001822A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/44
; CURRENT APPLICATION NUMBER: US/10/032,037B
; PRIOR FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 204
; LENGTH: 266
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-204

Query Match      100.0%; Score 29; DB 15; Length 266;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MEAPVI 6
   |||||
Db 121 MEAPVI 126

RESULT 6
US-10-029-988B-204
; Sequence 204, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; PRIOR FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 204
; LENGTH: 266
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-204

Query Match      100.0%; Score 29; DB 15; Length 266;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MEAPVI 6
   |||||
Db 121 MEAPVI 126

RESULT 7
US-10-032-423A-204
; Sequence 204, Application US/10032423A
; Publication No. US20040002450A1
; GENERAL INFORMATION:
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; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE OF INVENTION: MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/45
; CURRENT APPLICATION NUMBER: US/10/032,423A
; PRIOR FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 204
; LENGTH: 266
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-423A-204

Query Match      100.0%; Score 29; DB 15; Length 266;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 8
US-10-029-926B-25
; Sequence 25, Application US/10029926B
; Publication No. US20040073011A1
; GENERAL INFORMATION:
; APPLICANT: HAGAY, et al.
; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
; FILE REFERENCE: 10793/50
; CURRENT APPLICATION NUMBER: US/10/029,926B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 203
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 25
; LENGTH: 277
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-926B-25

Query Match      100.0%; Score 29; DB 12; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 9.
US-10-032-037B-25
; Sequence 25, Application US/10032037B
; Publication No. US20040001822A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE OF INVENTION: MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/44
; CURRENT APPLICATION NUMBER: US/10/032,037B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 25
; LENGTH: 277
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-25

Query Match      100.0%; Score 29; DB 15; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 10
US-10-029-988B-25
; Sequence 25, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE OF INVENTION: MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 25
; LENGTH: 277
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-25

Query Match      100.0%; Score 29; DB 15; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 11
US-10-032-423A-25
; Sequence 25, Application US/10032423A
; Publication No. US20040002450A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE OF INVENTION: MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/45
; CURRENT APPLICATION NUMBER: US/10/032,423A
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 25
; LENGTH: 277
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-423A-25

Query Match      100.0%; Score 29; DB 15; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 12
US-10-369-493-7871
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US-10-032-037B-25

Query Match      100.0%; Score 29; DB 15; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 10
US-10-029-988B-25
; Sequence 25, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE OF INVENTION: MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 25
; LENGTH: 277
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-25

Query Match      100.0%; Score 29; DB 15; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 11
US-10-032-423A-25
; Sequence 25, Application US/10032423A
; Publication No. US20040002450A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE OF INVENTION: MOETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/45
; CURRENT APPLICATION NUMBER: US/10/032,423A
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 25
; LENGTH: 277
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-423A-25

Query Match      100.0%; Score 29; DB 15; Length 277;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      121 MRAPVI 126

RESULT 12
US-10-369-493-7871
```

```
; Sequence 7871, Application US/10369493
; Publication No. US20030233675A1
; GENERAL INFORMATION:
; APPLICANT: Cao, Yongwei
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Goldman, Barry S.
; APPLICANT: Chen, Xianfeng
; TITLE OF INVENTION: EXPRESSION OF MICROBIAL PROTEINS IN PLANTS FOR PRODUCTION OF
; TITLE OF INVENTION: PLANTS WITH IMPROVED PROPERTIES
; FILE REFERENCE: 38-10(52052)B
; CURRENT APPLICATION NUMBER: US/10/369,493
; CURRENT FILING DATE: 2003-02-28
; PRIOR APPLICATION NUMBER: US 60/360,039
; PRIOR FILING DATE: 2002-02-21
; NUMBER OF SEQ ID NOS: 47374
; SEQ ID NO 7871
; LENGTH: 442
; TYPE: PRT
; ORGANISM: Rhodobacter sphaeroides
US-10-369-493-7871

Query Match      100.0%; Score 29; DB 15; Length 442;
Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      77 MRAPVI 82

RESULT 13
US-10-029-926B-26
; Sequence 26, Application US/10029926B
; Publication No. US20040073011A1
; GENERAL INFORMATION:
; APPLICANT: HAGAY, et al.
; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
; FILE REFERENCE: 10793/50
; CURRENT APPLICATION NUMBER: US/10/029,926B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 203
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 26
; LENGTH: 464
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-926B-26

Query Match      100.0%; Score 29; DB 12; Length 464;
Best Local Similarity 100.0%; Pred. No. 3.1e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      118 MRAPVI 123

RESULT 14
US-10-032-037B-26
; Sequence 26, Application US/10032037B
; Publication No. US20040001822A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/44
; CURRENT APPLICATION NUMBER: US/10/032,037B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
```

```
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 26
; LENGTH: 464
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-26

Query Match      100.0%; Score 29; DB 15; Length 464;
Best Local Similarity 100.0%; Pred. No. 3.1e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      118 MRAPVI 123

RESULT 15
US-10-029-988B-26
; Sequence 26, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 26
; LENGTH: 464
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-26

Query Match      100.0%; Score 29; DB 15; Length 464;
Best Local Similarity 100.0%; Pred. No. 3.1e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
Db      118 MRAPVI 123

Search completed: September 18, 2004, 04:20:41
Job time : 239.286 secs
```


GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:33:04 ; Search time 12.2143 Seconds
(without alignments)
47.252 Million cell updates/sec

Title: US-10-029-926B-8

Perfect score: 29

Sequence: 1 MRAPVI 6

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : PIR 78:*

1: pir1:*

2: pir2:*

3: pir3:*

4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	DB	ID	Description
1	29	100.0	249	2	H87675		hydrolase, alpha/b
2	29	100.0	442	2	JC4733		probable H-transp
3	29	100.0	629	2	T37255		acetylcholinestera
4	28	96.6	425	2	S33045		hypothetical prote
5	27	93.1	247	2	H69030		coenzyme PQQ synth
6	26	83.7	286	2	E85103		tagatase-bisphosph
7	26	89.7	286	2	A91131		tagatose-1,6-bisph
8	26	89.7	286	2	A85976		tagatose-bisphosph
9	26	89.7	327	2	E72057		ferrochelatase CP0
10	26	89.7	327	2	H86565		ferrochelatase [im
11	26	89.7	450	1	S13730		pmbA protein - Esc
12	26	89.7	450	2	D81280		maturatation of anti
13	26	89.7	450	2	D86121		maturatation of anti
14	26	89.7	450	2	A81056		probable PmbA prot
15	25	86.2	138	2	T24916		hypothetical prote
16	25	86.2	147	2	C95907		probable protein i
17	25	86.2	147	2	F95924		hypothetical prote
18	25	85.2	149	2	C72419		conserved hypothet
19	25	86.2	154	2	A92812		hypothetical prote
20	25	86.2	163	2	A87412		hypothetical prote
21	25	86.2	195	2	F75399		antibiotic resist
22	25	86.2	196	2	G75405		probable amidotran
23	25	86.2	203	2	C84409		imidazoleglycerol-
24	25	86.2	222	2	S56195		probable membrane
25	25	86.2	222	2	S63320		probable membrane
26	25	86.2	224	2	S85081		hypothetical prote
27	25	86.2	227	2	C53304		transfer protein C
28	25	86.2	240	2	E69004		hypothetical prote
29	25	86.2	252	2	A88508		protein H1412.4 i

30 25 86.2 261 2 AC3507 hisf protein [limpo
31 25 86.2 266 2 AC2407 tryptophan synthas
32 25 86.2 284 2 AD1712 fructose-1,6-bipho
33 25 86.2 284 2 AF1341 fructose-1,6-bipho
34 25 86.2 328 2 G96024 conserved hypothet
35 25 86.2 332 2 C81285 glyceraldehyde 3-p
36 25 86.2 359 2 AI0112 fructose-bisphosph
37 25 86.2 362 2 F90200 hypothetical prote
38 25 86.2 398 2 B70752 hypothetical prote
39 25 86.2 422 2 AE3394 lipoprotein releas
40 25 86.2 422 2 E83083 probable two-compo
41 25 86.2 430 2 T16715 hypothetical prote
42 25 86.2 446 2 A84940 pmbA protein [limpo
43 25 86.2 463 2 AC0977 L-seryl-tRNAse se
44 25 86.2 487 2 T27353 hypothetical prote
45 25 86.2 489 2 JC4787 shaw protein - Cal

ALIGNMENTS

RESULT 1

H87675
hydrolase, alpha/beta hydrolase fold family [imported] - Caulobacter crescentus
C;Species: Caulobacter crescentus
C;Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr-2001
C;Accession: H87675
B;Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.
B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolo
n, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A;Title: Complete Genome Sequence of Caulobacter crescentus.
A;Reference number: AB7249; MUID:21173698; PMID:11259647
A;Accession: H87675
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-249 <STO>
A;Cross-references: GB:AE005673; NID:gl3425158; PIDN:AAK25404.1; GSPDB:GNO0148
C;Genetics:
A;Gene: CC3442

Query Match 100.0%; Score 29; DB 2; Length 249;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 MRAPVI 6

Db 1 MRAPVI 6

RESULT 2

JC4733
probable H-transporing two-sector ATPase (EC 3.6.3.14), flagellum-specific - Rhodobac
C;Species: Rhodobacter sphaeroides
C;Date: 10-May-1996 #sequence_revision 19-Jul-1996 #text_change 03-Jun-2002
C;Accession: JC4733
R;Ballado, T.; Campos, A.; Camarena, L.; Dreyfus, G.
Gene 170, 69-72, 1996
A;Title: Flagellar genes from Rhodobacter sphaeroides are homologous to genes of the fl
A;Reference number: JC4733; MUID:96200857; PMID:8621091
A;Accession: JC4733
A;Molecule type: DNA
A;Residues: 1-442 <BAL>
A;Cross-references: GB:U31090; NID:gl518877; PIDN:AAB07344.1; PID:gl293928
C;Genetics:
A;Gene: fliI
C;Superfamily: H-transporing ATP synthase alpha chain; H-transporing ATP synthase a
C;Keywords: ATP; flagellum; hydrolase; nucleotide binding; P-loop
F;168-175/Region: nucleotide-binding motif A (P-loop)
F;191-362/Domain: H-transporing ATP synthase alpha chain homology <ATP>
F;191-207/Region: ATP-binding #status predicted
F;242-258/Domain: beta chain #status predicted <BET>

Query Match 100.0%; Score 29; DB 2; Length 442;
 Best Local Similarity 100.0%; Pred. No. 24;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||
 Db 77 MRAPVI 82

RESULT 3
 T37255
 acetylcholinesterase (EC 3.1.1.7) 2 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 21-Jul-2000
 C:Accession: T37255
 R:Grauso, M.; Culetto, E.; Combes, D.; Fedon, Y.; Toutant, J.P.; Arpagaus, M.
 F:BS Lett. 424, 279-284, 1998
 A:Title: Existence of four acetylcholinesterase genes in the nematodes Caenorhabditis el
 A:Reference number: Z21648; MUID:98198570; PMID:9539167
 A:Accession: T37255
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-629 <GRA>
 A:Cross-references: EMBL:AF025378; NID:G5149937; PIDN:AAC14016.2; PID:G5091489
 A:Experimental source: strain N2
 C:Genetics:
 A:Gene: ace-2
 A:Map position: I
 A:Superfamily: cholinesterase; cholinesterase homology
 C:Keywords: carboxylic ester hydrolase

Query Match 100.0%; Score 29; DB 2; Length 629;
 Best Local Similarity 100.0%; Pred. No. 34;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||
 Db 1 MRAPVI 6

RESULT 4
 S33045
 hypothetical protein - human herpesvirus 4
 C:Species: human herpesvirus 4, Epstein-Barr virus
 C:Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 20-Jun-2000
 C:Accession: S33045
 R:Farrell, P.J.
 submitted to the EMBL Data Library, March 1988
 A:Reference number: S32973
 A:Accession: S33045
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-425 <FAR>
 A:Cross-references: EMBL:V01555; NID:G59074; PIDN:CAA24796.1; PID:G1334904
 C:Superfamily: equine herpesvirus 2 hypothetical protein 23

Query Match 96.6%; Score 28; DB 2; Length 425;
 Best Local Similarity 83.3%; Pred. No. 41;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||
 Db 87 MRAPVI 92

RESULT 5
 H69030
 coenzyme PQQ synthetase protein III - Methanobacterium thermoautotrophicum (strain Delta
 C:Species: Methanobacterium thermoautotrophicum
 C:Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 22-Oct-1999
 C:Accession: H69030
 R:Smith, D.R.; Doucette-Stamm, L.A.; Deloughery, C.; Lee, H.; Dubois, J.; Aldredge, T.;
 Qiu, D.; Spadafora, R.; Vicaire, R.; Wang, Y.; Wierzbowski, J.; Gibson, R.; Jiawani, N.

ki, S.; Church, G.W.; Daniels, C.J.; Mao, J.; Rice, P.; Noelling, J.; Reeve, J.N.
 J. Bacteriol. 179, 7135-7155, 1997
 A:Title: Complete genome sequence of Methanobacterium thermoautotrophicum Delta H: funct
 A:Reference number: A69000; MUID:98037514; PMID:9371463
 A:Accession: H69030
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-247 <MTH>
 A:Cross-references: GB:AE000890; GB:AE000666; NID:G2622331; PIDN:AAB85716.1; PID:G2622333
 A:Experimental source: strain Delta H
 C:Genetics:
 A:Gene: MTH1227
 A:Start codon: GTG
 C:Keywords: iron; metalloprotein
 F:34, 38, 41/Binding site: iron (Cys) #status predicted
 Query Match 93.1%; Score 27; DB 2; Length 247;
 Best Local Similarity 66.7%; Pred. No. 42;
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||
 Db 4 MRAPVI 9

RESULT 6
 E65103
 tagatose-bisphosphate aldolase agay (EC 4.1.2.-) - Escherichia coli (strain K-12)
 C:Species: Escherichia coli
 C:Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 01-Mar-2002
 C:Accession: E65103
 R:Rattnet, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; Co
 A:Rose, D.J.; Mau, B.; Shao, Y.
 Science 277, 1453-1462, 1997
 A:Title: The complete genome sequence of Escherichia coli K-12.
 A:Reference number: A64720; MUID:97426617; PMID:9278503
 A:Accession: E65103
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-286 <BLAT>
 A:Cross-references: GB:AE000395; GB:U00096; NID:G1789524; PIDN:AAC76171.1; PID:G1789526;
 A:Experimental source: strain K-12, substrain MG1655
 C:Genetics:
 A:Gene: agay
 C:Superfamily: fructose-bisphosphate aldolase II
 C:Keywords: aldehyde-lyase; carbon-carbon lyase

Query Match 89.7%; Score 26; DB 2; Length 286;
 Best Local Similarity 83.3%; Pred. No. 87;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||
 Db 41 MRAPVI 46

RESULT 7
 A91131
 tagatose-1,6-bisphosphate aldolase [imported] - Escherichia coli (strain O157:H7, subst
 C:Species: Escherichia coli
 C:Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 03-Aug-2001
 C:Accession: A91131
 R:Havashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.
 Gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
 DNA Res. 8, 11-22, 2001
 A:Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and geno
 A:Reference number: A99629; MUID:21156231; PMID:11258796
 A:Accession: A91131
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-286 <HAY>
 A:Cross-references: GB:BA000007; PIDN:BA037440.1; PID:G13363490; GSPDB:GN00154
 A:Experimental source: strain O157:H7, substrain RMD 0509952

C:Genetics:

A:Gene: ECs4017

C:Superfamily: fructose-bisphosphate aldolase II

Query Match 89.7%; Score 26; DB 2; Length 286;

Best Local Similarity 83.3%; Pred. No. 87;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

Db 41 MRSPVI 46

RESULT 8

A85976

tagatose-bisphosphate aldolase 2 [imported] - Escherichia coli (strain O157:H7, substrain

C:Species: Escherichia coli

C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001

C:Accession: A85976

R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew

iller, L.; Grotbeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,

Nature 409, 529-533, 2001

A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.

A:Reference number: A85480; MUID:2107435; PMID:11206551

A:Accession: A85976

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-286 <STO>

A:Cross-references: GB:AE005174; NID:g12517735; PIDN:AAG58269.1; GSPDB:GN00145; UWGP:244

A:Experimental source: strain O157:H7, substrain EOL933

C:Genetics:

A:Gene: agay

C:Superfamily: fructose-bisphosphate aldolase II

Query Match 89.7%; Score 26; DB 2; Length 286;

Best Local Similarity 83.3%; Pred. No. 87;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

Db 41 MRSPVI 46

RESULT 9

E72057

ferrochelatase CP0144 [imported] - Chlamydia pneumoniae (strains CWL029 and AR39)

C:Species: Chlamydia pneumoniae, Chlamydia pneumoniae

C:Date: 23-Apr-1999 #sequence_revision 23-Apr-1999 #text_change 11-May-2000

C:Accession: E72057; F81609

R:Kelman, S.; Mitchell, W.; Marathe, R.; Lammel, C.; Fan, J.; Olinger, L.; Greenwood, J.

Nature Genet. 21, 385-389, 1999

A:Title: Comparative genomes of Chlamydia pneumoniae and C. trachomatis.

A:Reference number: A72000; MUID:99206606; PMID:10192388

A:Accession: E72057

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-327 <ARN>

A:Cross-references: GB:AE001645; GB:AE001363; NID:g4376896; PIDN:AA018742.1; PID:g437689

A:Experimental source: strain CWL029

R:Read, T.D.; Brunham, R.C.; Shen, C.; Gill, S.R.; Heidelberg, J.F.; White, O.; Hickey,

C.; Dodson, R.; Gwinn, M.; Nelson, W.; DeBoy, R.; Kolonay, J.; McClarty, G.; Salzberg,

Nucleic Acids Res. 28, 1397-1406, 2000

A:Title: Genome sequences of Chlamydia trachomatis MoPn and Chlamydia pneumoniae AR39.

A:Reference number: A81500; MUID:20150255; PMID:10684935

A:Accession: F81609

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-327 <REA>

A:Cross-references: GB:AE002175; GB:AE002161; NID:g7189069; PIDN:AAF38026.1; PID:g718907

A:Experimental source: strain AR39, HL cells

C:Genetics:

A:Gene: hem2; CP0144

C:Superfamily: ferrochelatase

Query Match 89.7%; Score 26; DB 2; Length 327;

Best Local Similarity 83.3%; Pred. No. 1e+02;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

Db 86 LRAPVI 91

RESULT 10

H86565

ferrochelatase [imported] - Chlamydia pneumoniae (strain J138)

C:Species: Chlamydia pneumoniae, Chlamydia pneumoniae

C:Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 02-Mar-2001

C:Accession: H86565

R:Shirai, M.; Hirakawa, H.; Kimoto, M.; Tabuchi, M.; Kishi, F.; Ouchi, K.; Shiba, T.;

Nucleic Acids Res. 28, 2311-2314, 2000

A:Title: Comparison of whole genome sequences of Chlamydia pneumoniae J138.

A:Reference number: A86491; MUID:20330349; PMID:10871362

A:Accession: H86565

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-327 <STO>

A:Cross-references: GB:BA000008; NID:g9978975; PIDN:BAA98810.1; GSPDB:GN00142

A:Experimental source: strain J138

C:Genetics:

A:Gene: hem2

Query Match 89.7%; Score 26; DB 2; Length 327;

Best Local Similarity 83.3%; Pred. No. 1e+02;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

Db 86 LRAPVI 91

RESULT 11

S13730

pmbA protein - Escherichia coli (strain K-12)

C:Species: Escherichia coli

C:Date: 29-Jan-1999 #sequence_revision 29-Jan-1999 #text_change 01-Mar-2002

C:Accession: S13730; S56461; F65235

R:Rodriguez-Sainz, M.C.; Hernandez-Chico, C.; Moreno, F.

Mol. Microbiol. 4, 1921-1932, 1990

A:Title: Molecular characterization of pmbA, an Escherichia coli chromosomal gene requi.

A:Reference number: S13730; MUID:91186828; PMID:2082149

A:Accession: S13730

A:Molecule type: DNA

A:Residues: 1-450 <ROD>

A:Cross-references: EMBL:X54152; NID:g42439; PIDN:CAA38091.1; PID:g42440

R:Burland, V.; Plunkett III, G.; Sofia, H.J.; Daniels, D.L.; Blattner, F.R.

Nucleic Acids Res. 23, 2105-2119, 1995

A:Title: Analysis of the Escherichia coli genome VI: DNA sequence of the region from 92

A:Reference number: S56314; MUID:95334362; PMID:7610040

A:Accession: S56461

A:Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-450 <BUR>

A:Cross-references: EMBL:U14003; NID:g1263172; PIDN:AAA97132.1; PID:g537077

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, August 1994

R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; C.

A.; Rose, D.J.; Mau, B.; Shao, Y.

Science 277, 1453-1462, 1997

A:Title: The complete genome sequence of Escherichia coli K-12.

A:Reference number: A64720; MUID:97426617; PMID:9278503

A:Accession: F65235

A:Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-450 <BLAT>

A:Cross-references: GB:AE000494; GB:U00096; NID:g1790670; PIDN:AACT7192.1; PID:g1790682

A:Experimental source: strain K-12, substrain MGL655

C;Genetics:

A;Gene: pmbA

A;Map position: 96 min

C;Superfamily: Escherichia coli pmbA protein

Query Match 89.7%; Score 26; DB 1; Length 450;

Best Local Similarity 83.3%; Pred. No. 1.4e+02;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

|:||||

Db 242 MKAPVI 247

RESULT 12

D91280

maturation of antibiotic MccB17 [imported] - Escherichia coli (strain O157:H7, substrain

C;Species: Escherichia coli

C;Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 03-Aug-2001

C;Accession: D91280

R;Hayashi, T.; Makino, K.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.

gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.

DNA Res. 8; 11-22, 2001

A;Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and gen

A;Reference number: A99629; MUID:21156231; PMID:11258796

A;Accession: D91280

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-450 <HAV>

A;Cross-references: GB:BA000007; PIDN:BA38635.1; PID:gl3364589; GSPDB:GN00154

A;Experimental source: strain O157:H7, substrain RMD 0509952

C;Genetics:

A;Gene: ECs212

C;Superfamily: Escherichia coli pmbA protein

Query Match

Best Local Similarity 89.7%; Score 26; DB 2; Length 450;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

|:||||

Db 242 MKAPVI 247

RESULT 13

D86121

maturation of antibiotic MccB17, see tld genes [imported] - Escherichia coli (strain O15

C;Species: Escherichia coli

C;Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 14-Sep-2001

C;Accession: D86121

R;Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew

Miller, L.; Grobeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,

Nature 409, 529-533, 2001

A;Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.

A;Reference number: A85480; MUID:21074935; PMID:11206551

A;Accession: D86121

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-450 <STO>

A;Cross-references: GB:AB005174; NID:gl21519236; PIDN:AAG59432.1; GSPDB:GN00145; UWGP:258

A;Experimental source: strain O157:H7, substrain EDL933

C;Genetics:

A;Gene: pmbA

C;Superfamily: Escherichia coli pmbA protein

Query Match

Best Local Similarity 89.7%; Score 26; DB 2; Length 450;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

|:||||

Db 242 MKAPVI 247

RESULT 14

AE1056

probable PmbA protein pmbA [imported] - Salmonella enterica subsp. enterica serovar Typhi

C;Species: Salmonella enterica subsp. enterica serovar Typhi

A;Note: this species has also been called Salmonella typhi

C;Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 18-Nov-2002

C;Accession: AE1056

R;Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; Wain, J.; Churcher,

th, T.; Conington, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.; White, N.; Farrar,

S.; Moule, S.; O'Gaora, P.

Nature 413, 848-852, 2001

A;Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.;

A;Title: Complete genome sequence of a multiple drug resistant Salmonella enterica serov

A;Reference number: AB0502; MUID:21534947; PMID:11677608

A;Accession: AE1056

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-450 <PAR>

A;Cross-references: GB:AL513382; PIDN:CAD06898.1; PID:gl6505546; GSPDB:GN00176

C;Genetics:

A;Gene: pmbA

C;Superfamily: Escherichia coli pmbA protein

Query Match

Best Local Similarity 89.7%; Score 26; DB 2; Length 450;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

|:||||

Db 242 MKAPVI 247

RESULT 15

T24916

hypothetical protein T14G10.4 - Caenorhabditis elegans

C;Species: Caenorhabditis elegans

C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999

C;Accession: T24916

R;Wild, A.

submitted to the EMBL Data Library, January 1996

A;Reference number: Z19954

A;Accession: T24916

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-138 <WIL>

A;Cross-references: EMBL:Z68880; PIDN:CAA93096.1; GSPDB:GN00022; CESP:T14G10.4

A;Experimental source: clone T14G10

C;Genetics:

A;Gene: CESP:T14G10.4

A;Map position: 4

A;Introns: 52/3; 108/2

Query Match

Best Local Similarity 86.2%; Score 25; DB 2; Length 138;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPV 5

|:||||

Db 1 MRAPV 5

Search completed: September 18, 2004, 03:47:30

JOB time : 15.2143 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 02:29:47 ; Search time 8.78571 Seconds
(without alignments)
35.560 Million cell updates/sec

Title: US-10-029-926B-8

Perfect score: 29

Sequence: 1 MRAPVI 6

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	96.6	164	1 RHBI_RAT	O88779 rattus norv
2	28	96.6	373	1 RHBI_MOUSE	Q8VC82 mus musculu
3	28	96.6	425	1 YRL1_EBV	P30119 Epstein-bar
4	28	96.6	438	1 RHBI_HUMAN	O75783 homo sapien
5	27	93.1	535	1 TIEL_ERARE	O31168 brachydanio
6	26	89.7	250	1 PXJ1_BRAVA	O69158 bradyrhizob
7	26	89.7	286	1 AGAY_ECOLI	P42908 escherichia
8	26	89.7	327	1 HEMZ_CHLPN	Q927V1 chlamydia p
9	26	89.7	450	1 PMBA_ECOLI	P24231 escherichia
10	25	86.2	95	1 CH10_XANAC	Q8rit8 xanthomonas
11	25	86.2	215	1 HIS5_STRAW	Q82aa2 streptomyc
12	25	86.2	222	1 YFG0_YEAST	P43544 saccharomyc
13	25	86.2	222	1 YN74_YEAST	P53823 saccharomyc
14	25	86.2	224	1 YMY5_YEAST	Q03144 saccharomyc
15	25	86.2	258	1 HIS6_RHIME	Q92tb3 rhizobium m
16	25	86.2	261	1 HIS6_BRUME	O8ve37 brucella me
17	25	86.2	261	1 HIS6_BRUSU	O8fy07 brucella su
18	25	86.2	398	1 Y106_MYCTU	Q10899 mycobacteri
19	25	86.2	446	1 PMBA_EUCAL	P57191 buchnera ap
20	25	86.2	463	1 SELA_SALTI	Q8z2d8 salmonella
21	25	86.2	463	1 SELA_SALTY	Q8z169 salmonella
22	25	86.2	486	1 COBB_STROO	Q9rj16 streptomyc
23	25	86.2	510	1 INO1_HORVU	O65195 hordeum vul
24	25	86.2	545	1 SYFB_METMA	Q8pta3 methanosarc
25	25	86.2	550	1 VGLE_HSV11	P04489 herpes simp
26	25	86.2	1220	1 IF2P_HUMAN	O60841 homo sapien
27	24	82.8	67	1 Y7K8_BPF22	P14108 bacterioph
28	24	82.8	109	1 YS88_CABEL	Q09384 caenorhabdi
29	24	82.8	122	1 INL3_MOUSE	O09107 mus musculu
30	24	82.8	162	1 HRS3_HUMAN	P33816 homo sapien
31	24	82.8	186	1 YG61_METJA	Q39055 methanococc
32	24	82.8	216	1 ALKB_ECOLI	P50500 escherichia
33	24	82.8	216	1 ALKB_SALTY	P37462 salmonella

34	24	82.8	243	1 TPIS_PROMM	Q7v7d2 prochloroco
35	24	82.8	243	1 TPIS_SYNPX	Q7u804 synechococc
36	24	82.8	248	1 TPIS_CLOAB	O52533 clostridium
37	24	82.8	248	1 TPIS_CLOPE	Q8xxu1 clostridium
38	24	82.8	252	1 AMPR_HUMAN	P15514 homo sapien
39	24	82.8	252	1 TPIS_LACPL	Q887h4 lactobacill
40	24	82.8	253	1 TPIS_STAAM	Q925c3 staphylococ
41	24	82.8	253	1 TPIS_STAEP	Q88cd5 staphylococ
42	24	82.8	259	1 IF2C_GALSU	O08910 galdieria s
43	24	82.8	263	1 HIS6_RHILO	Q98ct1 rhizobium l
44	24	82.8	287	1 NAPH_ECOLI	P33934 escherichia
45	24	82.8	323	1 SOPB_ECOLI	P08867 escherichia

ALIGNMENTS

RESULT 1
RHBI_RAT ID RHBI_RAT STANDARD; PRT; 164 AA.
AC O88779;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Rhomboid-related protein 1 (EC 3.4.21.-) (RRP) (Rhomboid-like protein
DE 1) (Fragment)
GN RHBDL1 OR RHBDL
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Intestinal epithelium;
RX MEDLINE=98324821; PubMed=9662444;
RA Pascall J.C., Brown K.D.;
RT "Characterization of a mammalian cDNA encoding a protein with high
RT sequence similarity to the Drosophila regulatory protein Rhomboid.";
RL FEBS Lett. 429:337-340(1998).
CC -1- FUNCTION: May be involved in regulated intramembrane proteolysis
CC and the subsequent release of functional polypeptides from their
CC membrane anchors (By similarity).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- SIMILARITY: Belongs to peptidase family S54.
CC
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CC
CC -----
CC EMBL; Y17258; CAA76716.1; -
CC MEROPS; S54.950; -
CC InterPro; IP002610; Peptidase_S54.
CC Pfam; PF01694; Rhomboid; 1
CC Hydrolase; Protease; Serine protease; Transmembrane.
CC NON_TER 1
CC TRANSMEM 10 30 POTENTIAL.
CC TRANSMEM 32 52 POTENTIAL.
CC TRANSMEM 56 76 POTENTIAL.
CC TRANSMEM 120 140 POTENTIAL.
CC ACT_SITE 60 60 CHARGE RELAY SYSTEM (BY SIMILARITY).
CC ACT_SITE 125 125 CHARGE RELAY SYSTEM (BY SIMILARITY).
CC NON_TER 164 164
CC SQ SEQUENCE 164 AA; 17662 MW; CF62ACE3B6C99210 CRC64;
Query Match 96.6%; Score 28; DB 1; Length 164;
Best Local Similarity 83.3%; Pred. No. 6.3;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6

```
Db
|||||:
52 MRAPV 57

RESULT 2
RHB1_MOUSE
ID RHB1_MOUSE STANDARD; PRT; 373 AA.
AC Q8VC82.
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Rhomboid-related protein 1 (EC 3.4.21.-) (RRP) (Rhomboid-like protein
1).
GN RHBDL1 OR RHBDL.
OS Mus musculus (Mouse)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6; TISSUE=Retina;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg K.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Haieh F.,
RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M.J., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
RA Rickards S., McGowan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Ketterman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.J., Skalska U., Smallos D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
CC -!- FUNCTION: May be involved in regulated intramembrane proteolysis
and the subsequent release of functional polypeptides from their
membrane anchors (By similarity).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein (Probable).
CC -!- SIMILARITY: Belongs to peptidase family S54.
CC
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CC
CC EMBL; BC021549; AAH21549.1; -.
DR MEROPS; S54.950; -.
DR MGD; MGI:2384891; Rbdl.
DR InterPro; IPR002048; EF-hand.
DR InterPro; IPR002610; Peptide_S54.
DR Pfam; PF01694; Rhomboid; 1.
KW Hydrolase; Protease; Serine protease; Transmembrane.
FT TRANSMEM 131 151 POTENTIAL.
FT TRANSMEM 196 216 POTENTIAL.
FT TRANSMEM 219 239 POTENTIAL.
FT TRANSMEM 243 263 POTENTIAL.
FT TRANSMEM 275 294 POTENTIAL.
FT TRANSMEM 307 327 POTENTIAL.
FT TRANSMEM 340 360 POTENTIAL.
FT ACT_SITE 199 199 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 247 247 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 312 312 CHARGE RELAY SYSTEM (BY SIMILARITY).

SQ SEQUENCE 373 AA; 41786 MW; 1F4E538B3A363D2A CRC64;
Query Match 96.6%; Score 28; DB 1; Length 373;
Best Local Similarity 83.3%; Pred. No. 15;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 239 MRAPV 244
|||||:

RESULT 3
YTRI_EBV
ID YTRI_EBV STANDARD; PRT; 425 AA.
AC P30L19.
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE Hypothetical BTRF1 protein.
GN BTRF1.
OS Epstein-barr virus (strain B95-8) (Human herpesvirus 4).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Gammaherpesvirinae; Lymphocryptovirus.
OX NCBI_TaxID=10377;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=84270667; PubMed=6087149;
RA Baer R., Bankier A.T., Biggin M.D., Deininger P.L., Farrell P.J.,
RA Gibson T.J., Hatfull G., Hudson G.S., Satchwell S.C., Seguin C.,
RA Tuffnell P.S., Barrett B.G.;
RA "DNA sequence and expression of the B95-8 Epstein-Barr virus genome.";
RL Nature 310:207-211 (1984).
CC -!- SIMILARITY: TO HVS-1 GENE 23.
CC
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CC
CC EMBL; V01555; CA24796.1; -.
DR EIR; S33045; S33045.
DR InterPro; IPR006772; Herpes_BTRF1.
DR Pfam; PF04682; Herpes_BTRF1; 1.
KW Hypothetical protein.
SQ SEQUENCE 425 AA; 46711 MW; 0ECCE5FD30495BD CRC64;
Query Match 96.6%; Score 28; DB 1; Length 425;
Best Local Similarity 83.3%; Pred. No. 18;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 87 MRAPV 92
|||||:

RESULT 4
RHB1_HUMAN
ID RHB1_HUMAN STANDARD; PRT; 438 AA.
AC Q75783; Q9NQ85;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Rhomboid-related protein 1 (EC 3.4.21.-) (RRP) (Rhomboid-like protein
1).
GN RHBDL1 OR RHBDL.
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
```

```
RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
RC TISSUE=Leukemia;
RX MEDLINE=98324821; PubMed=9662444;
RA Pascall J.C., Brown K.D.;
RT "Characterization of a mammalian cDNA encoding a protein with high
RL sequence similarity to the Drosophila regulatory protein Rhomboid.";
RN FEBS Lett. 429:337-340(1998).
[2]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RX MEDLINE=21096910; PubMed=11157797;
RA Daniels R.J., Peden J.F., Lloyd C., Horsley S.W., Clark K.,
RA Tufarelli C., Kearney L., Buckle V.J., Doggett N.A., Flint J.,
RA Higgs D.R.;
RT "Sequence, structure and pathology of the fully annotated terminal 2
RL Mb of the short arm of human chromosome 16.";
RN Hum. Mol. Genet. 10:339-352(2001).
CC -!- FUNCTION: May be involved in regulated intramembrane proteolysis
CC and the subsequent release of functional polypeptides from their
CC membrane anchors (By similarity).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1;
CC IsoId=O75783-1; Sequence=Displayed;
CC Name=2;
CC IsoId=O75783-2; Sequence=VSP_005372;
CC -!- TISSUE SPECIFICITY: Detected in heart, brain, skeletal muscle and
CC kidney.
CC -!- SIMILARITY: Belongs to peptidase family S54.
CC
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CC
CC EMBL; Y17108; CAA75629.1; -
CC EMBL; AJ272344; CAC00640.1; -
CC EMBL; AE006464; AAK61241.1; -
CC Genew; HGNC:10007; RHBDL1.
CC MIM; 603264; -
CC MEMOPS; S54.950; -
CC GO; GO:0005887; C:integral to plasma membrane; TAS.
CC GO; GO:0005624; C:membrane fraction; TAS.
CC GO; GO:0007165; P:signal transduction; TAS.
CC InterPro; IPR002048; EF-hand.
CC InterPro; IPR002610; Peptidase_S54.
CC Pfam; PF01694; Rhomboid; 1.
CC Hydrolase; Protease; Serine protease; Transmembrane;
KW Alternative splicing.
FT TRANSMEM 196 216 POTENTIAL.
FT TRANSMEM 262 282 POTENTIAL.
FT TRANSMEM 284 304 POTENTIAL.
FT TRANSMEM 308 328 POTENTIAL.
FT TRANSMEM 340 359 POTENTIAL.
FT TRANSMEM 372 392 POTENTIAL.
FT TRANSMEM 405 425 POTENTIAL.
FT ACT_SITE 264 264 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 312 312 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 377 377 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT VARSPLIC 1 77 MGRVEDGGTTEEDDPGTSALPAPGKQGPREGTGTGPL
FT SOKWEPPDAPSGQPGALMSRGARTQALAGGSSL -> M
FT DRSSLLQLIOR (in isoform 2).
FT /FTId=VSP_005372
SQ SEQUENCE 438 AA; 48314 MW; A7644AD3644A2P6 CRC64;
Query Match 96.6%; Score 28; DB 1; Length 438;
Best Local Similarity 83.3%; Pred. No. 18;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 MRAPV1 6
```

```
Db 304 MRAPV1 309
RESULT 5
TLE1_BRARE STANDARD; PRT; 535 AA.
ID TLE1_BRARE
AC O13188;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Groucho 1 protein (Fragment).
GN GRC1 OR GROUCHO1.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RA Wuelbeck C., Campos-Ortega J.A.;
RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: NUCLEAR EFFECTOR MOLECULE (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: Nuclear (Probable).
CC -!- SIMILARITY: Contains 6 WD repeats.
CC -!- SIMILARITY: Belongs to the WD-repeat Groucho/TLE family.
CC PROTEINS.
CC
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CC
CC EMBL; Y12467; CAA73070.1; -
CC EMBL; U96451; AAB57808.1; -
CC ZFIN; ZDB-GENE-990415-85; Grc1.
CC InterPro; IPR001680; WD40.
CC Pfam; PF00400; WD40; 6.
CC ProDom; PD000018; WD40; 1.
CC SMART; SM00320; WD40; 7.
CC PROSITE; PS00678; WD_REPEATS_1; 2.
CC PROSITE; PS00082; WD_REPEATS_2; 2.
CC PROSITE; PS00294; WD_REPEATS_REGION; 1.
KW Nuclear protein; Repeat; WD repeat.
FT NON_TER 1 1
FT DOMAIN <1 42 CCN DOMAIN.
FT DOMAIN 43 214 SER/PRO-RICH.
FT REPEAT 235 266 WD 1.
FT REPEAT 293 323 WD 2.
FT REPEAT 337 367 WD 3.
FT REPEAT 379 409 WD 4.
FT REPEAT 461 491 WD 5.
FT REPEAT 502 532 WD 6.
SQ SEQUENCE 535 AA; 57360 MW; A201A32071101363 CRC64;
Query Match 93.1%; Score 27; DB 1; Length 535;
Best Local Similarity 83.3%; Pred. No. 41;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 MRAPV1 6
Db 192 MRAPV1 197
RESULT 6
PDJX_BRAJA STANDARD; PRT; 250 AA.
ID PDJX_BRAJA
AC O69158;
DT 30-MAY-2000 (Rel. 39, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
```

10-OCT-2003 (Rel. 42, Last annotation update)
Pyridoxal phosphate biosynthetic protein pdxJ (PNP synthase).
PDXJ OR BL15064.
Bradyrhizobium japonicum.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Bradyrhizobiaceae; Bradyrhizobium.
OX NCBI_TaxID=375;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 1105PC4;
RA Mueller P., Stangel D.;
RT "Extended DNA sequencing in the upstream region of sipF in
Bradyrhizobium japonicum."
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiyama T.,
RA Sasamoto S., Watanabe A., Ideawa K., Iiguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimpo S., Tsurutaka H., Wada T., Yamada M.,
RA Tabata S.;
RT "Complete genomic sequence of nitrogen-fixing symbiotic bacterium
Bradyrhizobium japonicum USDA110."
RL DNA Res. 9:189-197(2002).
RN [3]
RP SEQUENCE OF 67-250 FROM N.A.
RC STRAIN=USDA 1105PC4;
RX MEDLINE=99086246; PubMed=9870699;
RA Baird A., Mueller P.;
RT "A second gene for type I signal peptidase in Bradyrhizobium
japonicum, sipF, is located near genes involved in RNA processing and
cell division."
RL Mol. Gen. Genet. 260:346-356(1998).
CC -!- FUNCTION: Catalyzes the condensation of 1-deoxy-D-xylulose-5-
phosphate (DXP) and L-amino-3-(phosphohydroxy)propan-2-one to form
pyridoxine 5'-phosphate (PNP) (By similarity).
CC -!- PATHWAY: De novo synthesis of pyridoxine (vitamin B6) and
pyridoxal phosphate.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (Probable).
CC -!- SIMILARITY: Belongs to the pdxJ family.
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CC -----
DR EMBL; AF065159; AAD02936.2; -;
DR EMBL; AP005953; BAC50329.1; -;
DR HSSP; P24223; 1H01.
DR HAMAP; MF_00279; -; 1.
DR InterPro; IPR004569; PdxJ.
DR Pfam; PF03740; PdxJ; 1.
DR ProDom; PD013657; PdxJ; 1.
DR TIGRFAMs; TIGR00559; pdxJ; 1.
KW Pyridoxine biosynthesis; Complete proteome.
FT CONFLICT 185 186
FT CONFLICT 238 240 SMR -> NA (IN REF. 1 AND 3).
FT CONFLICT 250 AA; 27033 MW; 47DBD939C0FA59CC CRC64;
SQ SEQUENCE 250 AA; 27033 MW; 47DBD939C0FA59CC CRC64;
Query Match 89.7%; Score 26; DB 1; Length 250;
Best Local Similarity 83.3%; Pred. NO. 33;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 WRAPVI 6
Db 149 LRAPVI 154
RESULT 7

AGAY_ECOLI
ID AGAY_ECOLI STANDARD; PRT; 286 AA.
AC P42908;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Tagatose-1,6-bisphosphate aldolase agay (EC 4.1.2.-) (TBPA).
GN AGAY OR KBA OR B3137 OR C3894 OR Z4491 OR ECS4017.
OS Escherichia coli,
OS Escherichia coli O6, and
OS Escherichia coli O157:H7.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=562, 217992, 83334;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;
RA Blattner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12."
RL Science 277:1453-1474(1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=O6:H1 / CFT073 / ATCC 700928;
RX MEDLINE=22388234; PubMed=12471157;
RA Welch R.A., Burland V., Plunkett G. III, Redford P., Roesch P.,
RA Rasko D., Buckles E.L., Liou S.-R., Boutin A., Hackett J., Stroud D.,
RA Mayhew G.F., Rose D.J., Zhou S., Schwartz D.C., Perna N.T.,
RA Mobley H.L.T., Donnenberg M.S., Blattner F.R.;
RT "Extensive mosaic structure revealed by the complete genome sequence
of uropathogenic Escherichia coli."
RL Proc. Natl. Acad. Sci. U.S.A. 99:17020-17024(2002).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / EDL933 / ATCC 700927;
RX MEDLINE=21074935; PubMed=11206551;
RA Perna N.T., Plunkett G. III, Burland V., Mau B., Glasner J.D.,
RA Rose D.J., Mayhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,
RA Posfai G., Hackett J., Klink S., Boutin A., Shao Y., Miller L.,
RA Grobeck E.J., Davis N.W., Lim A., Dimalanta E.T., Potamousis K.,
RA Apodaca J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,
RA Welch R.A., Blattner F.R.;
RT "Genome sequence of enterohaemorrhagic Escherichia coli O157:H7."
RL Nature 409:529-533(2001).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / RIMD 0509552;
RX MEDLINE=21156231; PubMed=11258796;
RA Hayashi T., Makino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,
RA Han C.-G., Ohtsubo E., Nakayama K., Murata T., Tanaka M., Tobe T.,
RA Iida T., Takami H., Honda T., Sasakawa C., Ogasawara N., Yasunaga T.,
RA Kuhara S., Shiba T., Hattori M., Shinagawa H.;
RT "Complete genome sequence of enterohaemorrhagic Escherichia coli
O157:H7 and genomic comparison with a laboratory strain K-12."
RL DNA Res. 8:11-22(2001).
RN [5]
RP DISCUSSION OF SEQUENCE.
RX MEDLINE=97086503; PubMed=8932697;
RA Reizer J., Ramseier T.M., Reizer A., Charbit A., Saier M.H. Jr.;
RT "Novel phosphotransferase genes revealed by bacterial genome
sequencing: a gene cluster encoding a putative N-acetylglucosamine
metabolic pathway in Escherichia coli."
RL Microbiology 142:231-250(1996).
RN [6]
RP X-RAY CRYSTALLOGRAPHY (1.45 ANGSTROMS).
RX MEDLINE=22050650; PubMed=11940603;
RA Hall D.R., Bond C.S., Leonard G.A., Watt C.I., Berry A., Hunter W.N.;
RT "Structure of tagatose-1,6-bisphosphate aldolase. Insight into chiral
discrimination, mechanism, and specificity of class II aldolases."
RL J. Biol. Chem. 277:22018-22024(2002).

DT 01-MAR-1992 (Rel. 21, Created)
 DT 01-MAR-1992 (Rel. 21, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE PMBA protein (Tlde protein).
 GN PMBA OR TLDE OR B4235 OR Z5845 OR ECS5212.
 OS Escherichia coli, and
 OS Escherichia coli O157:H7.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
 OC Enterobacteriaceae; Escherichia.
 OX NCBI_TaxID=562, 83334;
 RN [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=K12;
 RC MEDLINE=91186828; PubMed=2082149;
 RA Rodriguez-Sainz M.C., Hernandez-Chico C., Moreno F.,
 RT "Molecular characterization of pmBA, an Escherichia coli chromosomal
 gene required for the production of the antibiotic peptide MccB17.";
 RL Mol. Microbiol. 4:1921-1932(1990).
 RN [2]
 RN SEQUENCE FROM N.A., AND CHARACTERIZATION.
 RC STRAIN=K12;
 RC MEDLINE=96177756; PubMed=8604133;
 RA Murayama N., Shimizu H., Takiguchi S., Baba Y., Amino H.,
 RA Horiuchi T., Sekimizu K., Miki T.;
 RT "Evidence for involvement of Escherichia coli genes pmBA, csrA and a
 previously unrecognized gene tldD, in the control of DNA gyrase by
 RT letB (cddB) of sex factor F.";
 RL J. Mol. Biol. 256:483-502(1996).
 RN [3]
 RN SEQUENCE FROM N.A.
 RC STRAIN=K12 / MG1655;
 RC MEDLINE=95334362; PubMed=7610040;
 RA Burland V.D., Plunkett G. III, Sofia H.J., Daniels D.L.,
 RA Blattner F.R.;
 RT "Analysis of the Escherichia coli genome VI: DNA sequence of the
 RT region from 92.8 through 100 minutes.";
 RL Nucleic Acids Res. 23:2105-2119(1995).
 RN [4]
 RN SEQUENCE FROM N.A.
 RC STRAIN=O157:H7 / EDL933 / ATCC 700927;
 RC MEDLINE=21074935; PubMed=11206551;
 RA Perna N.T., Plunkett G. III, Burland V., Mau B., Glasner J.D.,
 RA Rose D.J., Mayhew G.P., Evans P.S., Gregor J., Kirkpatrick H.A.,
 RA Postfai G., Hackett J., Klink S., Boutin A., Shao Y., Miller L.,
 RA Grobeck E.J., Davis N.W., Lim A., Dimalanta E.T., Potamocis K.,
 RA Apodaca J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,
 RA Welch R.A., Blattner F.R.;
 RT "Genome sequence of enterohaemorrhagic Escherichia coli O157:H7.";
 RL Nature 409:529-533(2001).
 RN [5]
 RN SEQUENCE FROM N.A.
 RC STRAIN=O157:H7 / RIMD 0509952;
 RC MEDLINE=21156231; PubMed=11258796;
 RA Hayashi T., Makino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,
 RA Han C.-G., Ohtsubo E., Nakayama K., Murata T., Tanaka M., Tobe T.,
 RA Iida T., Takami H., Honda T., Sasaki C., Ogasawara N., Yasunaga T.,
 RA Kuhara S., Shiba T., Hattori M., Shinagawa H.;
 RT "Complete genome sequence of enterohaemorrhagic Escherichia coli
 RT O157:H7 and genomic comparison with a laboratory strain K-12.";
 RL DNA Res. 8:11-22(2001).
 CC -!- FUNCTION: MAY FACILITATE THE SECRETION OF THE PEPTIDE ANTIBIOTIC
 MICROBIN B17 (MCCB17) BY COMPLETING ITS MATURATION. SUPPRESSES THE
 INHIBITORY ACTIVITY OF THE CARBON STORAGE REGULATOR (CSRA).
 CC -!- SUBCELLULAR LOCATION: Cytoplasmic.
 CC -!- SIMILARITY: Belongs to the tldD/pmbA family.
 CC
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CC
 DR EMBL; X54152; CAA38091.1; -
 DR EMBL; D4452; BAA07915.1; -
 DR EMBL; U14003; AAA97132.1; -
 DR EMBL; AB000494; AAC77192.1; -
 DR EMBL; AB005655; AAG59432.1; -
 DR EMBL; AP002568; BAB38635.1; -
 DR PIR; D86121; D86121.
 DR PIR; D91280; D91280.
 DR PIR; S13730; S13730.
 DR EcoGene; EGI0741; pmBA.
 DR InterPro; IPR002510; Peptidase_U62.
 DR Pfam; PF01523; PmbA_Tlde; 1.
 KW Complete proteome.
 SQ SEQUENCE 450 AA; 48369 MW; A062969197B52B5E CRC64;
 Query Match 89.7%; Score 26; DB 1; Length 450;
 Best Local Similarity 83.3%; Pred. No. 62;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MEAPVI 6
 DB 242 MKAPVI 247
 ID CH10_XANAC STANDARD; PRT; 95 AA.
 AC Q8RTG;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE 10 kDa chaperonin (Protein Cpn10) (groES protein).
 GN GROS OR GROS OR XAC0541 OR XCC0522.
 OS Xanthomonas axonopodis (pv. citri).
 OS Xanthomonas campestris (pv. campestris), and
 OS Xanthomonas campestris (pv. phaseoli).
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
 OC Xanthomonadaceae; Xanthomonas.
 OX NCBI_TaxID=92829, 340, 29445;
 RN [1]
 RN SEQUENCE FROM N.A.
 RC SPECIES=X.a. citri, and X.c. campestris;
 RC STRAIN=306 / ATCC 13902 / XY 101, and ATCC 33913 / NCPPB 528;
 RC MEDLINE=22022445; PubMed=12024217;
 RA da Silva A.C.R., Ferro J.A., Reinach F.C., Parah C.S., Furlan L.R.,
 RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
 RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.B.A.,
 RA Camarotte G., Cannavan F., Cardozo J., Chambergo F., Ciapina L.P.,
 RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorry H.,
 RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Perro M.I.T.,
 RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
 RA Katayama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
 RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
 RA Martins E.C., Medeiros J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
 RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
 RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
 RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
 RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
 RA Setubal J.C., Kitajima J.P.;
 RT "Comparison of the genomes of two Xanthomonas pathogens with differing
 RT host specificities.";
 RL Nature 417:459-463(2002).
 RN [2]
 RN SEQUENCE FROM N.A.
 RC SPECIES=X.c. phaseoli;
 RA Hsu C.-C., Yu Y.-J., Yang M.-T.;
 RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Binds to Cpn60 in the presence of Mg-ATP and suppresses
 the ATPase activity of the latter.
 CC -!- SUBUNIT: Heptamer of 7 subunits arranged in a ring (By
 similarity).
 CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).

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CC  -!- SIMILARITY: Belongs to the groES chaperonin family.
CC -----
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CC -----
CC EMBL; AE011680; AAM35430.1; -
CC EMBL; AE012149; AAM39838.1; -
CC EMBL; AF426387; AAL74149.1; -
CC HAMAP; MF_00860; -; 1.
CC InterPro; IPR001476; Chaprinin_Cpn10.
CC Pfam; PF00166; cpn10; 1.
CC PRINTS; PR00297; CHAPERONIN10.
CC ProDom; PD000566; Chaprinin_Cpn10; 1.
CC PROSITE; PS00681; CHAPERONINS_CPN10; 1.
CC Chaperone; Complete proteome.
CC SEQUENCE 95 AA; 9979 MW; POF0724BD153CB3 CRC64;
CC -----
CC Query Match 86.2%; Score 25; DB 1; Length 95;
CC Best Local Similarity 66.7%; Pred. No. 21;
CC Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
CC -----
CC QY 1 MRAPVI 6
CC Db 54 LRAPVV 59
CC -----
CC RESULT 11
CC HIS5_STRAW STANDARD; PRT; 215 AA.
CC ID Q82AA2;
CC DT 15-MAR-2004 (Rel. 43, Created)
CC DT 15-MAR-2004 (Rel. 43, Last sequence update)
CC DE Imidazole glycerol phosphate synthase subunit hisH (EC 2.4.2.-) (IGP
CC synthase glutamine amidotransferase subunit) (IGP synthase subunit
CC hisH) (ImGP synthase subunit hisH) (IGPS subunit hisH).
CC GN HIS5 OR SAV6157.
CC OS Streptomyces avermitilis.
CC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
CC Streptomycinae; Streptomycetaceae; Streptomycetes.
CC NCBI_TaxID=33903;
CC RN [1]
CC SEQUENCE FROM N.A.
CC RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
CC EX MEDLINE=21477403; PubMed=11572948;
CC RA Omura S., Ikeda H., Ishikawa J., Hanamoto A., Takahashi C.,
CC Shinose M., Takahashi Y., Horikawa H., Nakazawa H., Osonoe T.,
CC Kikuchi H., Shiba T., Sakaki Y., Hattori M.;
CC RT "Genome sequence of an industrial microorganism Streptomyces
CC avermitilis: deducing the ability of producing secondary
CC metabolites."
CC RT Proc. Natl. Acad. Sci. U.S.A. 98:12215-12220(2001).
CC RN [2]
CC SEQUENCE FROM N.A.
CC RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
CC RX MEDLINE=22609306; PubMed=12692562;
CC RA Ikeda H., Ishikawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba T.,
CC Sakaki Y., Hattori M., Omura S.;
CC RT "Complete genome sequence and comparative analysis of the industrial
CC microorganism Streptomyces avermitilis."
CC RL Nat. Biotechnol. 21:526-531(2003).
CC -!- FUNCTION: IGP synthase catalyzes the conversion of PRFAR and glutamine to
CC IGP, AICAR and glutamate. The hisH subunit provides the glutamine
CC amidotransferase activity that produces the ammonia necessary to
CC hisF for the synthesis of IGP and AICAR (By similarity).
CC -!- CATALYTIC ACTIVITY: 5-[(5-phospho-1-deoxybulos-1-
CC ylaminomethylideneamino)-1-(5-phosphoribosyl)imidazole-4-
CC carboxamide + L-glutamine = imidazole-glycerol phosphate + 5-

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CC aminoimidazol-4-carboxamide ribonucleotide + L-glutamate + H(2)O.
CC -!- PATHWAY: Histidine biosynthesis; fifth step.
CC -!- SUBUNIT: Heterodimer of hisH and hisF (By similarity).
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Contains 1 type-1 glutamine amidotransferase domain.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF005045; BAC73868.1; -
CC HAMAP; MF_00278; -; 1.
CC InterPro; IPR000991; GATase_1.
CC Pfam; PF00117; GATase; 1.
CC PROSITE; PS00442; GATASE_TYPE_I; 1.
CC Histidine biosynthesis; Transferase; Glutamine amidotransferase;
CC Complete proteome.
CC FT ACT_SITE 86 86 BY SIMILARITY.
CC FT ACT_SITE 196 196 BY SIMILARITY.
CC FT ACT_SITE 198 198 BY SIMILARITY.
CC SQ SEQUENCE 215 AA; 23052 MW; DDAD5CC85943CBBE CRC64;
CC -----
CC Query Match 86.2%; Score 25; DB 1; Length 215;
CC Best Local Similarity 66.7%; Pred. No. 51;
CC Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
CC -----
CC QY 1 MRAPVI 6
CC Db 167 MRAPLV 172
CC -----
CC RESULT 12
CC YFG0_YEAST STANDARD; PRT; 222 AA.
CC ID P435Z4;
CC DT 01-NOV-1995 (Rel. 32, Created)
CC DT 01-NOV-1995 (Rel. 32, Last sequence update)
CC DT 15-MAR-2004 (Rel. 43, Last annotation update)
CC DE Hypothetical protein YFL060C.
CC GN YFL060C.
CC OS Saccharomyces cerevisiae (Baker's Yeast).
CC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
CC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
CC NCBI_TaxID=4932;
CC RN [1]
CC SEQUENCE FROM N.A.
CC RC STRAIN=S288C / AB972;
CC RX MEDLINE=95400292; PubMed=7670463;
CC RA Murakami Y., Naitou M., Hagiwara H., Shibata T., Ozawa M.,
CC Sasanuma S.-I., Sasanuma M., Tsuchiya Y., Soeda E., Yokoyama K.,
CC Yamazaki M., Tashiro H., Eki T.;
CC RT "Analysis of the nucleotide sequence of chromosome VI from
CC Saccharomyces cerevisiae."
CC RL Nat. Genet. 10:261-268(1995).
CC -!- SIMILARITY: Belongs to the UPF0030 family.
CC -----
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CC -----
CC EMBL; D50617; BAA09181.1; -
CC PIR; S56195; S56195.
CC GerMOnline; 140096; -
CC SGD; S0001834; YFL060C.
CC GO; GO:000515; F:protein binding; IPI.

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DR GO:0008614; P:pyridoxine metabolism; IDA.
DR GO:0009228; P:thiamin biosynthesis; IMP.
DR InterPro: IPR002161; SNO.
DR Pfam: PF01174; SNO; 1.
DR PROSITE: PS01236; UPF0030; 1.
KW Hypothetical protein.
SQ SEQUENCE 222 AA; 25132 MW; 6755EE1D826CAABA CRC64;

Query Match      86.2%; Score 25; DB 1; Length 222;
Best Local Similarity 83.3%; Pred. No. 53;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
DB      151 IRAPVI 156

RESULT 13
VN74_YEAST
ID VN74_YEAST STANDARD; PRT; 222 AA.
AC PS3823;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Hypothetical UPF0030 protein YNL334C.
GN YNL334C OR N0285.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97313269; PubMed=9169873;
RA Philippsen P., Kleine K., Pohlmann R., Duesterhoft A., Hamberg K.,
RA Hegmann J.H., Obermaier B., Urrestazu L.A., Aert R., Albermann K.,
RA Altmann R., Andre B., Baladron V., Ballesta J.P.G., Becan A.-M.,
RA Beinbauer J., Boskovic J., Butrago M.J., Bussereau F., Coster F.,
RA Crouzet M., D'Angelo M., Dal Pero F., De Antoni A., Del Rey F.,
RA Dolgoust M., Dondy H., Dubois E., Fiedler T., Fleig U., Floeth M.,
RA Fritz C., Gallardin C., Garcia-Cantalejo J.M., Glandsdorff N.,
RA Goffeau A., Guelender U., Herbert C.D., Heumann K., Heuss-Neitzel D.,
RA Hilbert H., Hinni K., Iraqui Housaini I., Jaquet M., Jimenez A.,
RA Joniaux J.-L., Karpfinger-Hartl L., Lanfranchi G., Lepingle A.,
RA Levesque H., Lyck R., Maftahi M., Mallet L., Maurer K.C.T.,
RA Messenguy F., Mewes H.-W., Moestl D., Nasr F., Nicaud J.-M.,
RA Niedenthal R.K., Pandolfo D., Pierard A., Piravandi E., Planta R.J.,
RA Pohl T.M., Purnelle B., Rebischung C., Remacha M., Revuelta J.L.,
RA Rinke M., Saiz J.E., Sartorello F., Scherens B., Sen-Gupta M.,
RA Soler-Mira A., Urbanus J.H.M., Valle G., Van Dyck L., Verhasselt P.,
RA Vierendeels F., Visiers S., Voet M., Volckaert G., Wach A.,
RA Wambutt R., Wedler K., Zollner A., Hani J.;
RT "The nucleotide sequence of Saccharomyces cerevisiae chromosome XIV
and its evolutionary implications.";
RL Nature 387:93-98(1997).
CC -!- SIMILARITY: Belongs to the UPF0030 family.
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; 271610; CAA96268.1; -.
CC PIR; S63320; S63320.
CC GenOnline; 143340; -.
CC SGD; S0005278; YNL334C.
CC GO; GO:0008614; P:pyridoxine metabolism; IDA.
CC GO; GO:0009228; P:thiamin biosynthesis; IMP.
CC InterPro; IPR002161; SNO.
CC Pfam; PF01174; SNO; 1.

RESULT 14
YMY5_YEAST
ID YMY5_YEAST STANDARD; PRT; 224 AA.
AC Q03144;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Hypothetical UPF0030 protein YMR095C.
GN YMR095C OR YM6543.02C.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97313268; PubMed=9169872;
RA Bowman S., Churcher C.M., Badcock K., Brown D., Chillingworth T.,
RA Connor R., Dedman K., Devlin K., Gentles S., Hamlin N., Hunt S.,
RA Jagals K., Lye G., Moule S., Odell C., Pearson D., Rajandream M.A.,
RA Rice P., Skelton J., Walsh S., Whitehead S., Barrett B.G.;
RT "The nucleotide sequence of Saccharomyces cerevisiae chromosome
XIII.";
RL Nature 387:90-93(1997).
CC -!- SIMILARITY: Belongs to the UPF0030 family.
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; 249807; CAA89896.1; -.
CC PIR; S55081; S55081.
CC GenOnline; 142763; -.
CC SGD; S0004701; YMR095C.
CC GO; GO:0008614; P:pyridoxine metabolism; IDA.
CC GO; GO:0009228; P:thiamin biosynthesis; IMP.
CC InterPro; IPR002161; SNO.
CC Pfam; PF01174; SNO; 1.
CC PROSITE; PS01236; UPF0030; 1.
KW Hypothetical protein.
SQ SEQUENCE 224 AA; 24906 MW; C0EC1B2382581D40 CRC64;

Query Match      86.2%; Score 25; DB 1; Length 224;
Best Local Similarity 83.3%; Pred. No. 53;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
DB      160 IRAPVI 165

RESULT 15
HIS6_RHIME
ID HIS6_RHIME STANDARD; PRT; 258 AA.
AC Q92T53;
DR 28-FEB-2003 (Rel. 41, Created)

```

```

DT 28-FEB-2003 (Rel. 41, Last sequence update)
DE 28-FEB-2003 (Rel. 41, Last annotation update)
DE Imidazole glycerol phosphate synthase subunit hisF (EC 4.1.3.-) (IGP
DE synthase cyclase subunit) (IGP synthase subunit hisF) (imGP synthase
DE subunit hisF) (IGPS subunit hisF).
GN HISF OR R00051 OR SMC02569.
OS Rhizobium meliloti (Sinorhizobium meliloti).
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Rhizobiaceae; Sinorhizobium/Ensifer group; Sinorhizobium.
OX NCBI_TaxID=382;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1021;
RX MEDLINE=21396507; PubMed=11481430;
RA Capela D., Barloy-Hubler F., Gouzy J., Bothe G., Ampe F., Batut J.,
RA Boistard P., Becker A., Boutry M., Cadieu E., Dreano S., Gloux S.,
RA Goudie T., Goffeau A., Kahn D., Kiss E., Lelaure V., Masuy D.,
RA Pohl T., Portetelle D., Puehler A., Purnelle B., Ramsperger U.,
RA Renard C., Thebaud P., Vandenbol M., Weidner S., Galibert F.;
RT "Analysis of the chromosome sequence of the legume symbiont
RT Sinorhizobium meliloti strain 1021."
RL Proc. Natl. Acad. Sci. U.S.A. 98:9877-9882(2001).
CC -!- FUNCTION: IGP synthase catalyzes the conversion of PRFAR and glutamine to
CC IGP, AICAR and glutamate. The hisF subunit catalyzes the
CC cyclization activity that produces IGP and AICAR from PRFAR using
CC the ammonia provided by the hisH subunit (By similarity).
CC -!- CATALYTIC ACTIVITY: 5-[(5-phospho-1-deoxyribulose-1-
CC ylamino)methylideneamino]-1-(5-phosphoribosyl)imidazole-4-
CC carboxamide + L-glutamine = imidazole-glycerol phosphate + 5-
CC aminoimidazol-4-carboxamide ribonucleotide + L-glutamate + H(2)O.
CC -!- PATHWAY: Histidine biosynthesis; fifth step.
CC -!- SUBUNIT: Heterodimer of hisH and hisF (By similarity).
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the hisA / hisF family.
CC -----
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CC -----
DR EMBL; AL591782; CAC41438.1; -.
DR HAMAP; MF_01013; -.
DR InterPro; IPR003009; FMN_enzyme.
DR InterPro; IPR006062; His_biosynth.
DR InterPro; IPR004651; HisF.
DR Pfam; PF00977; His_biosynth; 1.
DR TIGRFAMs; TIGR00735; hisF; 1.
KW Histidine biosynthesis; Lyase; Complete proteome.
FT ACT SITE 12 12 POTENTIAL.
FT ACT SITE 131 131 POTENTIAL.
SQ SEQUENCE 258 AA; 27312 MW; 56125BFB22F39D3 CRC64;
Query Match 86.2%; Score 25; DB 1; Length 258;
Best Local Similarity 83.3%; Pred. No. 62;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Qy 1 MRAPVI 6
Db 200 VRAPVI 205

```

Search completed: September 18, 2004, 03:43:44
Job time : 11.7857 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:24:30 ; Search time 31.2857 Seconds
(without alignments)
60.510 Million cell updates/sec

Title: US-10-029-926B-8

Perfect score: 29

Sequence: 1 MRAPVI 6

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

- 1: SP archaea:*
- 2: SP bacteria:*
- 3: SP fungi:*
- 4: SP human:*
- 5: SP invertebrate:*
- 6: SP mammal:*
- 7: SP mbc:*
- 8: SP organelle:*
- 9: SP phage:*
- 10: SP plant:*
- 11: SP rodent:*
- 12: SP virus:*
- 13: SP vertebrate:*
- 14: SP unclassified:*
- 15: SP virus:*
- 16: SP bacteriap:*
- 17: SP archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	29	100.0	249	16 Q9A2W4	Q9A2W4 caulobacter
2	29	100.0	442	2 Q53093	Q53093 rhodobacter
3	29	100.0	442	2 Q53153	Q53153 rhodobacter
4	29	100.0	629	5 Q61371	Q61371 caenorhabdi
5	29	100.0	947	11 Q80TL6	Q80TL6 mus musculu
6	28	96.6	376	16 Q8P9F5	Q8P9F5 xanthomonas
7	28	96.6	541	16 Q8NT69	Q8NT69 corynebacte
8	28	96.6	713	5 Q7YXV3	Q7YXV3 cryptospori
9	27	93.1	247	17 Q27235	Q27235 methanobact
10	27	93.1	376	2 Q9L778	Q9L778 xanthomonas
11	27	93.1	376	16 Q8PL69	Q8PL69 xanthomonas
12	27	93.1	416	16 Q9ACQ3	Q9ACQ3 streptomyce
13	27	93.1	449	16 Q89X81	Q89X81 bradyrhizob
14	27	93.1	771	13 Q98TH2	Q98TH2 brachydanio
15	26	89.7	162	5 Q8ILB7	Q8ILB7 plasmodium
16	26	89.7	286	2 Q9KIP8	Q9KIP8 escherichia

17	26	89.7	353	8 Q9TE71	Q9TE71 cylindrothe
18	26	89.7	353	8 Q9TE68	Q9TE68 nitzschia f
19	26	89.7	450	16 Q8FAP2	Q8FAP2 escherichia
20	26	89.7	450	16 Q8XG77	Q8XG77 salmorella
21	26	89.7	450	16 Q83P54	Q83P54 shigella fl
22	26	89.7	450	16 Q7UAN7	Q7UAN7 shigella fl
23	26	89.7	463	16 Q7UKY8	Q7UKY8 rhodopirell
24	26	89.7	535	16 Q8ERE8	Q8ERE8 oceanobacil
25	26	89.7	583	5 Q9VKES	Q9VKES drosophila
26	26	89.7	1127	13 Q7TIR3	Q7TIR3 brachydanio
27	26	89.7	1449	5 Q9V917	Q9V917 drosophila
28	25	86.2	93	12 Q8QNH1	Q8QNH1 ectocarpus
29	25	86.2	94	2 P97101	P97101 alcaligenes
30	25	86.2	103	16 Q83913	Q83913 enterococcu
31	25	86.2	110	10 Q9LJ06	Q9LJ06 cryza sativ
32	25	86.2	119	10 Q7XMD2	Q7XMD2 cryza sativ
33	25	86.2	138	5 Q22499	Q22499 caenorhabdi
34	25	86.2	147	2 Q9X699	Q9X699 rhizobium m
35	25	86.2	147	16 Q92W24	Q92W24 rhizobium m
36	25	86.2	147	16 Q92VN7	Q92VN7 rhizobium m
37	25	86.2	149	16 Q9WX16	Q9WX16 thermotoga
38	25	86.2	154	16 Q9FG98	Q9FG98 xylella fas
39	25	86.2	154	16 Q87A23	Q87A23 xylella fas
40	25	86.2	159	16 Q893T7	Q893T7 clostridium
41	25	86.2	163	16 Q9A8P1	Q9A8P1 caulobacter
42	25	86.2	182	16 Q8EDN5	Q8EDN5 shewanella
43	25	86.2	188	1 Q8UWY2	Q8UWY2 sulfobolus
44	25	86.2	188	16 Q7UWU0	Q7UWU0 rhodopirell
45	25	86.2	195	16 Q9RUG7	Q9RUG7 deinococcus

ALIGNMENTS

RESULT 1

Q9A2W4 ID Q9A2W4 PRELIMINARY; PRT; 249 AA.
AC Q9A2W4;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hydrolase, alpha/beta hydrolase fold family.
GN CC3442.
OS Caulobacter crescentus.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacteriales;
OC Caulobacteraceae; Caulobacter.
OX NCBI_TaxID=155892;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 19089 / CB15;
RX MEDLINE=21173698; PubMed=11259647;
RA Nierman W.C., Feldblyum T.V., Laub M.T., Paulsen I.T., Nelson K.E.,
Eisen J., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R.,
Potocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B.,
DeBoy R.T., Dodson R.J., Durkin A.S., Gwinn M.L., Haft D.H.,
Kolony J.F., Smit J., Craven M.B., Knouri H., Haft D.H., Berry K.,
Uterback T., Tran K., Wolf A., Vamathevan J., Shetty J., Berry K.,
Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;
RT "Complete genome sequence of Caulobacter crescentus.";
RL Proc Natl Acad. Sci. U.S.A. 98:4136-4141(2001).
CC -1- SIMILARITY: TO ALPHA/BETA HYDROLASE FOLD.
EMBL; AB006003; AK25404.1; -.
DR PIR; H87675; H87675.
DR TIGR; CC3442; -.
DR GO; GO:0016787; F:hydrolase activity; IEA.
DR GO; GO:0006725; P:aromatic compound metabolism; IEA.
DR InterPro; IPR000073; A/b hydrolase.
DR InterPro; IPR003089; AB_Hydrolase.
DR InterPro; IPR000379; Ser_estrs.
DR Pfam; PF00561; abhydrolase; 1.
DR PRINTS; PR00111; ABHYDROLASE.
KW Hydrolase; Complete proteome.
SQ SEQUENCE 249 AA; 26772 MW; 491350127DD300AE CRC64;

Query Match 100.0%; Score 29; DB 16; Length 249;
 Best Local Similarity 100.0%; Pred. No. 84;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OS Rhodobacter sphaeroides (Rhodospseudomonas sphaeroides).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhodobacterales;
 OC Rhodobacteraceae; Rhodobacter.
 RN [1] NCBI_TaxID=1063;
 RP SEQUENCE FROM N.A.
 RC STRAIN=WS8;
 RX MEDLINE=96327148; PubMed=8759796;
 RA Goodfellow I.G., Pollitt C.E., Sockett R.E.;
 RT "Cloning of the flII gene from Rhodobacter sphaeroides WS8 by analysis
 of a transposon mutant with impaired motility.";
 RL FEMS Microbiol. Lett. 142:111-116(1996).
 CC -|- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
 DR EMBL; X37201; CAA65834.1; -.
 DR GO; GO:0005737; C:cytoplasm; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0000166; F:nucleotide binding; IEA.
 DR GO; GO:0000058; P:biogenesis; IEA.
 DR GO; GO:0006811; P:ion transport; IEA.
 DR GO; GO:0015031; P:protein transport; IEA.
 DR GO; GO:0015992; P:proton transport; IEA.
 DR InterPro; IPR003593; AAA ATPase.
 DR InterPro; IPR00194; ATPase a/bcentre.
 DR InterPro; IPR005714; FliI_YscN.
 DR Pfam; PF00006; ATP-synt_ab; 1.
 DR SMART; SM00382; AAA; 1.
 DR TIGRFAWS; TIGR01026; fliI_yscN; 1.
 DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
 KW ATP-binding; Hydrogen ion transport; Hydrolase; Ion transport;
 KW Transport.
 SQ SEQUENCE 442 AA; 46813 MW; AD070D4E17FD3CC3 CRC64;

Query Match 100.0%; Score 29; DB 2; Length 442;
 Best Local Similarity 100.0%; Pred. No. 1.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OS Rhodobacter sphaeroides (Rhodospseudomonas sphaeroides).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhodobacterales;
 OC Rhodobacteraceae; Rhodobacter.
 RN [1] NCBI_TaxID=1063;
 RP SEQUENCE FROM N.A.
 RC STRAIN=WS8;
 RX MEDLINE=96327148; PubMed=8759796;
 RA Goodfellow I.G., Pollitt C.E., Sockett R.E.;
 RT "Cloning of the flII gene from Rhodobacter sphaeroides WS8 by analysis
 of a transposon mutant with impaired motility.";
 RL FEMS Microbiol. Lett. 142:111-116(1996).
 CC -|- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
 DR EMBL; X37201; CAA65834.1; -.
 DR GO; GO:0005737; C:cytoplasm; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0000166; F:nucleotide binding; IEA.
 DR GO; GO:0000058; P:biogenesis; IEA.
 DR GO; GO:0006811; P:ion transport; IEA.
 DR GO; GO:0015031; P:protein transport; IEA.
 DR GO; GO:0015992; P:proton transport; IEA.
 DR InterPro; IPR003593; AAA ATPase.
 DR InterPro; IPR00194; ATPase a/bcentre.
 DR InterPro; IPR005714; FliI_YscN.
 DR Pfam; PF00006; ATP-synt_ab; 1.
 DR SMART; SM00382; AAA; 1.
 DR TIGRFAWS; TIGR01026; fliI_yscN; 1.
 DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
 KW ATP-binding; Hydrogen ion transport; Hydrolase; Ion transport;
 KW Transport.
 SQ SEQUENCE 442 AA; 46852 MW; E6D35531F5A59BAE CRC64;

OS Rhodobacter sphaeroides (Rhodospseudomonas sphaeroides).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhodobacterales;
 OC Rhodobacteraceae; Rhodobacter.
 RN [1] NCBI_TaxID=1063;
 RP SEQUENCE FROM N.A.
 RC STRAIN=WS8;
 RX MEDLINE=96327148; PubMed=8759796;
 RA Goodfellow I.G., Pollitt C.E., Sockett R.E.;
 RT "Cloning of the flII gene from Rhodobacter sphaeroides WS8 by analysis
 of a transposon mutant with impaired motility.";
 RL FEMS Microbiol. Lett. 142:111-116(1996).
 CC -|- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
 DR EMBL; X37201; CAA65834.1; -.
 DR GO; GO:0005737; C:cytoplasm; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0000166; F:nucleotide binding; IEA.
 DR GO; GO:0000058; P:biogenesis; IEA.
 DR GO; GO:0006811; P:ion transport; IEA.
 DR GO; GO:0015031; P:protein transport; IEA.
 DR GO; GO:0015992; P:proton transport; IEA.
 DR InterPro; IPR003593; AAA ATPase.
 DR InterPro; IPR00194; ATPase a/bcentre.
 DR InterPro; IPR005714; FliI_YscN.
 DR Pfam; PF00006; ATP-synt_ab; 1.
 DR SMART; SM00382; AAA; 1.
 DR TIGRFAWS; TIGR01026; fliI_yscN; 1.
 DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
 KW ATP-binding; Hydrogen ion transport; Hydrolase; Ion transport;
 KW Transport.
 SQ SEQUENCE 442 AA; 46813 MW; AD070D4E17FD3CC3 CRC64;

Query Match 100.0%; Score 29; DB 2; Length 442;
 Best Local Similarity 100.0%; Pred. No. 1.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OS Rhodobacter sphaeroides (Rhodospseudomonas sphaeroides).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhodobacterales;
 OC Rhodobacteraceae; Rhodobacter.
 RN [1] NCBI_TaxID=1063;
 RP SEQUENCE FROM N.A.
 RC STRAIN=WS8;
 RX MEDLINE=96327148; PubMed=8759796;
 RA Goodfellow I.G., Pollitt C.E., Sockett R.E.;
 RT "Cloning of the flII gene from Rhodobacter sphaeroides WS8 by analysis
 of a transposon mutant with impaired motility.";
 RL FEMS Microbiol. Lett. 142:111-116(1996).
 CC -|- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
 DR EMBL; X37201; CAA65834.1; -.
 DR GO; GO:0005737; C:cytoplasm; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0015078; F:hydrogen ion transporter activity; IEA.
 DR GO; GO:0000166; F:nucleotide binding; IEA.
 DR GO; GO:0000058; P:biogenesis; IEA.
 DR GO; GO:0006811; P:ion transport; IEA.
 DR GO; GO:0015031; P:protein transport; IEA.
 DR GO; GO:0015992; P:proton transport; IEA.
 DR InterPro; IPR003593; AAA ATPase.
 DR InterPro; IPR00194; ATPase a/bcentre.
 DR InterPro; IPR005714; FliI_YscN.
 DR Pfam; PF00006; ATP-synt_ab; 1.
 DR SMART; SM00382; AAA; 1.
 DR TIGRFAWS; TIGR01026; fliI_yscN; 1.
 DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
 KW ATP-binding; Hydrogen ion transport; Hydrolase; Ion transport;
 KW Transport.
 SQ SEQUENCE 442 AA; 46813 MW; AD070D4E17FD3CC3 CRC64;

Query Match 100.0%; Score 29; DB 2; Length 442;
 Best Local Similarity 100.0%; Pred. No. 1.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RA None;
RT "Genome sequence of the nematode C. elegans: a platform for
investigating biology. The C. elegans Sequencing Consortium.";
RL Science 262:2012-2018 (1998).
[4]
RN SEQUENCE FROM N.A.
RC STRAIN=Bristol N2;
RA Woessner J., Graves T., Keppler D.;
RT "The sequence of C. elegans cosmid Y44E3A.";
RL Submitted (NOV-1998) to the EMBL/GenBank/DBJ databases.
[5]
RN SEQUENCE FROM N.A.
RC STRAIN=Bristol N2;
RA Waterston R.;
RT "Direct Submission.";
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO THE TYPE-B CARBOXYL ESTERASE/LIPASE FAMILY.
DR EMBL; AF025378; AAC14016.2; -;
DR EMBL; AF065899; AAC78228.2; -;
DR PIR; T33842; T33842.
DR PIR; T37255; T37255.
DR HSSP; P21836; 1VAA.
DR WormPep; Y44E3A.2; CE28363.
DR GO; GO:0003930; F:acetylcholinesterase activity; IEA.
DR GO; GO:0004104; F:cholinesterase activity; IEA.
DR GO; GO:0016787; F:hydrolase activity; IEA.
DR InterPro; IPR002018; Carboxesterases.
DR InterPro; IPR000997; Cholinesterase.
DR InterPro; IPR000379; Ser. esters.
DR Pfam; PF00135; Coesterase; 1.
DR PRINTS; P00878; CHOLNSTRASE.
DR PROSITE; PS00122; CARBOXYL ESTERASE_B_1; 1.
KW Hydrolase; Hypochemical protein.
SQ SEQUENCE 629 AA; 70863 MW; 74940F512FEDF869 CRC64;

Query Match 100.0%; Score 29; DB 5; Length 629;
Best Local Similarity 100.0%; Pred. No. 2.1e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
DB 1 MRAPVI 6

RESULT 5
Q80TL6 PRELIMINARY; PRT; 947 AA.
AC Q80TL6;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE MKIAA1042 protein (fragment).
GN MKIAA1042.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC Tissue=Brain;
RX MEDLINE=22579291; PubMed=12693553;
RA Okazaki N., Kikuno R., Ohara R., Inamoto S., Aizawa H., Yuasa S.,
RA Nakajima D., Nagase T., Ohara O., Koga H.;
RT "Prediction of the coding sequences of mouse homologues of KIAA gene:
II. The complete nucleotide sequences of 400 mouse KIAA-homologous
RT cDNAs identified by screening of terminal sequences of cDNA clones
randomly sampled from size-fractionated libraries.";
RL DNA Res. 10:35-48 (2003).
DR EMBL; AK122426; BAC65708.1; -;
DR GO; GO:0005634; C:nucleus; ISS.
DR GO; GO:0005886; C:plasma membrane; ISS.
DR GO; GO:0005515; F:protein binding; ISS.
DR GO; GO:0005102; F:receptor binding; ISS.

DR GO; GO:0006493; P:O-linked glycosylation; ISS.
DR GO; GO:0006605; P:protein targeting; ISS.
DR GO; GO:0006357; P:regulation of transcription from Pol II promoter; ISS.
DR InterPro; IPR006933; HAP1_N.
DR InterPro; IPR000585; Hemopexin.
DR Pfam; PF04849; HAP1_N; 1.
DR PROSITE; PS00024; HEMOPEXIN; 1.
FT NON_TER
SQ SEQUENCE 947 AA; 105486 MW; 0987284C6ACF223A5 CRC64;

Query Match 100.0%; Score 29; DB 11; Length 947;
Best Local Similarity 100.0%; Pred. No. 3.2e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
DB 924 MRAPVI 929

RESULT 6
Q8P9F5 PRELIMINARY; PRT; 376 AA.
AC Q8P9F5;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Flagellar protein.
GN FLHB OR XCC1910.
OS Xanthomonas campestris (pv. campestris).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xanthomonas.
OX NCBI_TaxID=340;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=ATCC 33913 / NCPPB 528;
RC MEDLINE=22022145; PubMed=12024217;
RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,
RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
RA Camarotte G., Camnava F., Cardozo J., Chambergo F., Ciapina L.P.,
RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorry H.,
RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Gruber A.,
RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
RA Trindade dos Santos M., Truffi D., Teal S.M., White F.F.,
RA Setubal J.C., Kitajima J.P.;
RT "Comparison of the genomes of two Xanthomonas pathogens with differing
RT host specificities.";
RL Nature 417:459-463 (2002).
DR EMBL; AB012294; AAM41199.1; -;
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0009306; P:protein secretion; IEA.
DR InterPro; IPR006135; Bac Export_2.
DR InterPro; IPR006136; FLHB.
DR Pfam; PF01312; Bac export 2; 1.
DR PRINTS; P00950; TYPE3MSPROT.
DR TIGRFAMs; TIGR00328; FLHB; 1.
KW Complete proteome.
SQ SEQUENCE 376 AA; 41490 MW; 406778385A158910 CRC64;

Query Match 96.6%; Score 28; DB 16; Length 376;
Best Local Similarity 83.3%; Pred. No. 2.2e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
DB 285 MRAPVI 290

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RESULT 7
Q8NT69 PRELIMINARY; PRT; 541 AA.
ID Q8NT69 AC
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE ResB protein required for cytochrome c biosynthesis.
GN CGL0441.
OS Corynebacterium glutamicum (Brevibacterium flavum).
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Corynebacteriaceae; Corynebacterium.
NCBI_TaxID=1718;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RA Nakagawa S.;
RT "Complete genomic sequence of Corynebacterium glutamicum ATCC 13032.";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBSJ databases.
DR EMBL; AF005275; BAB97834.1; -.
DR InterPro; IPR007816; ResB.
DR Pfam; PF05140; ResB; 1.
KW Complete proteome.
SQ SEQUENCE 541 AA; 61244 MW; 37F42BA0A74F78BB CRC64;

Query Match 96.6%; Score 28; DB 16; Length 541;
Best Local Similarity 83.3%; Pred. No. 3.2e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 103 MRAPVI 108

RESULT 8
Q7YVY3 PRELIMINARY; PRT; 713 AA.
ID Q7YVY3 AC
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN IMB.818.
OS Cryptosporidium parvum.
OC Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;
OC Cryptosporidiidae; Cryptosporidium.
OX NCBI_TaxID=5807;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Iowa;
RA Bankier A.T., Spriggs H.F., Fartmann B., Konfortov B.A., Madera M.,
RA Vogel C., Teichmann S.A., Ivens A., Dear P.H.;
RT "Integrated mapping, chromosomal sequencing and sequence analysis of
RT Cryptosporidium parvum.";
RL Genome Res. 0:0-0(2003).
DR EMBL; BX538353; CAD98347.1; -.
KW Hypothetical protein.
SQ SEQUENCE 713 AA; 82980 MW; 101BE54E3C2163CA CRC64;

Query Match 96.6%; Score 28; DB 5; Length 713;
Best Local Similarity 83.3%; Pred. No. 4.2e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 106 MRAPVI 111

RESULT 9
O27295 PRELIMINARY; PRT; 247 AA.
ID O27295 AC

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DT 01-JAN-1998 (TrEMBLrel. 05, Created)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Coenzyme PQQ synthesis protein III.
GN MTH1227.
OS Methanobacterium thermoautotrophicum.
OC Archaea; Euryarchaeota; Methanobacteria; Methanobacteriales;
OC Methanobacteriaceae; Methanothermobacter.
OX NCBI_TaxID=187420;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Delta H;
RX MEDLINE=98037514; PubMed=9371463;
RA Smith D.R., Doucette-Stamm L.A., DeLoughery C., Lee H.-M., Dubois J.,
RA Aldredge T., Bashirzadeh R., Blakely D., Cook R., Gilbert K.,
RA Harrison D., Hoang L., Keagle P., Lum W., Pothier B., Qiu D.,
RA Spadafora R., Vicare R., Wang Y., Wierzbowski J., Gibson R.,
RA Jiwani N., Caruso A., Bush D., Safer H., Patwell D., Prabhakar S.,
RA McDougall S., Shimer G., Goyal A., Pietrovski S., Church G.M.,
RA Daniels C.J., Mao J.-I., Rice P., Noelling J., Reeve J.N.;
RT "Complete genome sequence of Methanobacterium thermoautotrophicum
RT deltaH: functional analysis and comparative genomics.";
RL J. Bacteriol. 179:7135-7155(1997).
DR EMBL; AF000890; AAB85716.1; -.
DR PIR; H69030; H69030.
DR InterPro; IPR007197; Radical SAM.
DR Pfam; PF04055; Radical_SAM; 1.
KW Complete proteome.
SQ SEQUENCE 247 AA; 27374 MW; DD1F0C0251DC5217 CRC64;

Query Match 93.1%; Score 27; DB 17; Length 247;
Best Local Similarity 66.7%; Pred. No. 2.5e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
Db 4 MRAPVI 9

RESULT 10
Q9L778 PRELIMINARY; PRT; 376 AA.
ID Q9L778 AC
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Flagellar biosynthetic protein FlhB.
GN FLHB.
OS Xanthomonas oryzae (pv. oryzae).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xanthomonas.
OX NCBI_TaxID=64187;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21071238; PubMed=11204784;
RA Shen Y., Chern M.-S., Silva F.G., Ronald P.;
RT "Isolation of a Xanthomonas oryzae pv. oryzae flagellar operon region
RT and molecular characterization of flhB.";
RL Mol. Plant Microbe Interact. 14:204-213(2001).
DR EMBL; AF226282; AAF33828.1; -.
DR GO; GO:0046020; C:membrane; IEA.
DR GO; GO:0009306; P:protein secretion; IEA.
DR InterPro; IPR006135; Bac_Export_2.
DR InterPro; IPR006136; FlhB.
DR Pfam; PF01312; Bac export 2; 1.
DR PRINTS; PR00950; TYPE3IMSPROT.
DR TIGRPFAMs; TIGR00328; flhB; 1.
SQ SEQUENCE 376 AA; 41154 MW; 68514424C3D1E452 CRC64;

Query Match 93.1%; Score 27; DB 2; Length 376;
Best Local Similarity 66.7%; Pred. No. 3.8e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

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QY      1 MRAPVI 6
DB      285 MRAPIV 290

RESULT 11
Q8PL69 PRELIMINARY; PRT; 376 AA.
ID Q8PL69
AC Q8PL69;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Flagellar protein.
GN FLHB OR XAC1937.
OS Xanthomonas axonopodis (pv. citri).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xanthomonas.
OX NCBI_TaxID=92829;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=305 / ATCC 13902 / XV 1011;
RX MEDLINE=2202145; PubMed=12024217;
RA da Silva A.C.R., Ferro J.A., Reilbach F.C., Farah C.S., Furlan L.R.,
RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
RA Camarotte G., Cannavan F., Cardoso J., Chambergo F., Ciapina L.P.,
RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorri H.,
RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
RA Pereira H.A., Rossi A., Sena J.A.D., Silva C. de Souza R.F.,
RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Terza R.I.D.,
RA Trindade dos Santos M., Truffi D., Teal S.M., White F.F.,
RA Setubal J.C., Kitajima J.P.;
RT "Comparison of the genomes of two Xanthomonas pathogens with differing
RT host specificities."
RL Nature 417:459-463(2002).
DR EMBL; AE011823; AAC36799.1; -
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0009305; P:protein secretion; IEA.
DR InterPro; IPR006135; Bac Export_2.
DR InterPro; IPR006136; FlhB.
DR Pfam; PF01312; Bac export 2; 1.
DR PRINTS; PR00950; TYPE3INSEBOT.
DR TIGRFAMs; TIGR00328; flhB; 1.
KW Complete proteome.
SQ SEQUENCE 376 AA; 41333 MW; 615AE986871936AC CRC64;

Query Match 93.1%; Score 27; DB 16; Length 376;
Best Local Similarity 66.7%; Pred. No. 3.8e+02;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
DB      285 MRAPIV 290

RESULT 12
Q9ACQ9 PRELIMINARY; PRT; 416 AA.
ID Q9ACQ9
AC Q9ACQ9;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Putative transposon transposase.
GN SCPI.259.
OS Streptomyces coelicolor.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Plasmid SCPI.
OX NCBI_TaxID=92829;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiumi T.,
RA Sasamoto S., Watanabe A., Idesawa K., Iriguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimpo S., Tsuruoka H., Wada T., Yamada M.,
RA Tabata S.;

Query Match 93.1%; Score 27; DB 16; Length 416;
Best Local Similarity 83.3%; Pred. No. 4.2e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 MRAPVI 6
DB      232 MRAPVL 237

RESULT 13
Q89X81 PRELIMINARY; PRT; 449 AA.
ID Q89X81
AC Q89X81;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE BIR0433 protein.
GN BLR0433.
OS Bradyrhizobium japonicum.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Bradyrhizobiaceae; Bradyrhizobium.
OX NCBI_TaxID=375;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiumi T.,
RA Sasamoto S., Watanabe A., Idesawa K., Iriguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimpo S., Tsuruoka H., Wada T., Yamada M.,
RA Tabata S.;

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RT "Complete genomic sequence of nitrogen-fixing symbiotic bacterium
 RL Bradyrhizobium japonicum USDA110.";
 RL DNA Res. 9:189-197(2002).
 DR EMBL; AF005936; BAC45698.1; -;
 DR GO; GO:0004222; F:metalloendopeptidase activity; IEA.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro; IPR002886; Peptidase M37.
 DR Pfam; PF01551; Peptidase_M37; 1.
 KW Complete proteome.
 SQ SEQUENCE 449 AA; 47625 MW; 0374885B47AB3511 CRC64;

Query Match 93.1%; Score 27; DB 13; Length 449;
 Best Local Similarity 83.3%; Pred. No. 4.6e+02;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||:
 DB 1 MRAPVL 6

RESULT 14

Q98TH2 PRELIMINARY; PRT; 771 AA.
 AC Q98TH2;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-WAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Groucho-related gene 3 protein.
 GN GRG3 OR GRG3.
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OX NCBI_TaxID=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21294966; PubMed=11401394;
 RA Kobayashi M., Nishikawa K., Suzuki T., Yamamoto M.;
 RT "The homeobox protein Six3 interacts with the Groucho corepressor and
 acts as a transcriptional repressor in eye and forebrain formation.";
 RL Dev. Biol. 232:315-326(2001).
 DR EMBL; AB040707; BAB40698.1; -;
 DR ZFIN; ZDB-GENE-010410-1; Gr03.
 DR InterPro; IPR005617; TLE_N.
 DR InterPro; IPR001680; WD40.
 DR Pfam; PF03920; TLE_N; 1.
 DR Pfam; PF00400; WD40; 6.
 DR ProDom; PD000018; WD40; 1.
 DR SMART; SM00320; WD40; 7.
 DR PROSITE; PS00678; WD_REPEATS_1; 2.
 DR PROSITE; PS00082; WD_REPEATS_2; 2.
 DR PROSITE; PS0294; WD_REPEATS_REGION; 2.
 KW Repeat; WD repeat.
 SQ SEQUENCE 771 AA; 83789 MW; 3B9CEB203826E458 CRC64;

Query Match 93.1%; Score 27; DB 13; Length 771;
 Best Local Similarity 83.3%; Pred. No. 7.8e+02;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||:
 DB 428 MRAPVL 433

RESULT 15

Q8ILB7 PRELIMINARY; PRT; 162 AA.
 AC Q8ILB7;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Mitochondrial import inner membrane translocase subunit timl7,
 putative.

GN PF14_0328.
 OS Plasmodium falciparum (isolate 3D7).
 OC Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
 CX NCBI_TaxID=36329;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=3D7;
 RX MEDLINE=22255705; PubMed=12368864;
 RA Gardner M.J., Hall N., Fung E., White O., Barriman M., Hyman R.W.,
 RA Carlton J.M., Pain A., Nelson K.E., Bowman S., Paulsen I.T., James K.,
 RA Eisen J.A., Rutherford K., Salzberg S.L., Craig A., Kyes S.,
 RA Chan M.-S., Nene V., Shallow S.J., Suh B., Peterson J., Angiuoli S.,
 RA Partea M., Allen J., Selengut J., Haft D., Mather M.W., Vaidya A.B.,
 RA Martin D.M.A., Fairlamb A.H., Fraunholz M.J., Roos D.S., Ralph S.A.,
 RA McFadden G.I., Cummings L.M., Subramanian G.M., Mungall C., Venter J.C., Carucci D.J., Hoffman S.L., Newbold C., Davis R.W.,
 RA Fraser C.M., Barrell B.;
 RT "Genome sequence of the human malaria parasite Plasmodium falciparum.";
 RL Nature 419:498-511(2002).
 DR EMBL; AB014821; AAN36941.1; -;
 DR GO; GO:0005744; C:mitochondrial inner membrane pre-sequence t...; IEA.
 DR GO; GO:0015450; P:protein translocase activity; IEA.
 DR GO; GO:0015031; P:protein transport; IEA.
 DR InterPro; IPR003397; Tim17_Tim22.
 DR Pfam; PF02466; Tim17; 1.
 SQ SEQUENCE 162 AA; 17909 MW; F536704A462221FF CRC64;
 Query Match 89.7%; Score 26; DB 5; Length 162;
 Best Local Similarity 66.7%; Pred. No. 2.9e+02;
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRAPVI 6
 |||||:
 DB 58 MRAPIL 63

Search completed: September 18, 2004, 03:46:23
 Job time : 35.2857 secs

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OM protein - protein search, using sw model

Run on: September 18, 2004, 02:28:24; Search time 35.3571 Seconds
(without alignments)
39.956 Million cell updates/sec

Title: US-10-029-926b-114

Perfect score: 28

Sequence: 1 DYGMS 5

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: A_Geneseq_23Jan04.*
1: Geneseqp1980s.*
2: Geneseqp1990s.*
3: Geneseqp2000s.*
4: Geneseqp2001s.*
5: Geneseqp2002s.*
6: Geneseqp2003as.*
7: Geneseqp2003bs.*
8: Geneseqp2004s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	5	3 AAY95191	Aay95191 Anti-plat
2	28	100.0	5	5 ABG78239	Abg78239 Human Fv
3	28	100.0	5	5 ABG91930	Abg91930 Human ant
4	28	100.0	5	6 ABU11271	Abu11271 Human TAN
5	28	100.0	16	2 AAR49460	Aar49460 Factor VI
6	28	100.0	20	5 ABJ05002	Abj05002 A3 peptid
7	28	100.0	26	1 AAP61435	Aap61435 Factor VI
8	28	100.0	29	1 AAP50314	Aap50314 Peptide e
9	28	100.0	33	1 AAP50318	Aap50318 Peptide e
10	28	100.0	41	1 AAP50313	Aap50313 Thrombin
11	28	100.0	60	5 ABQ04948	Abq04948 A3 domain
12	28	100.0	73	3 AAB07206	Aab07206 Human Fac
13	28	100.0	98	3 AAB40073	Aab40073 Anti-H11
14	28	100.0	98	5 ABG78186	Abg78186 Human Fv
15	28	100.0	98	5 ABG91877	Abg91877 Human ant
16	28	100.0	98	6 ABQ27087	Abq27087 Human ger
17	28	100.0	113	3 AAY95177	Aay95177 Anti-plat
18	28	100.0	113	3 AAY95178	Aay95178 Anti-plat
19	28	100.0	115	3 AAY95189	Aay95189 Anti-plat
20	28	100.0	115	3 AAY95190	Aay95190 Anti-plat
21	28	100.0	116	2 AAM19880	Aam19880 CEA-speci
22	28	100.0	117	2 AAR66312	Aar66312 Human imm
23	28	100.0	118	4 AAU02560	Aau02560 Anti-adip
24	28	100.0	120	2 AAR25204	Aar25204 OP-G2 mon
25	28	100.0	207	5 AAU98019	Aau98019 Human ace

ALIGNMENTS

RESULT 1

AAY95191
ID AAY95191 standard; peptide; 5 AA.

XX AAY95191;

XX 29-AUG-2000 (first entry)

DE Anti-platelet glycoprotein Ib human H1b-1 VH CDR1.

XX Variable heavy chain; single chain antibody; scFv; human; H1b-1;
XX Glycoprotein Ib alpha; platelet; aggregation; antiaggregant;
XX antithrombotic; thrombus; therapy; diagnostic; CDR1;
XX Complementarity determining region.

OS Homo sapiens.

XX WO200026667-A1.

XX 11-MAY-2000.

XX 29-OCT-1999; 99WO-US025495.

XX 30-OCT-1998; 98US-0106275P.

XX (MILL/) MILLER J L.

XX Miller JL;

XX WPI; 2000-365744/31.

PT Isolated nucleic acid molecule encoding anti-human platelet glycoprotein
PT Ib alpha molecule useful for producing antibodies which inhibit platelet
PT aggregation.

XX Claim 14; Fig 5; 89pp; English.

XX The present sequence is that of complementarity determining region 1
XX (CDR1) of the heavy chain variable region (VH) of human single chain
XX antibody (scFv) H1b-1 (see AAY95198), which is directed against platelet
XX glycoprotein Ib (GPIb). The H1b series of scFv was isolated from a human
XX synthetic VH and VL scFv library on the basis of their binding to
XX platelet GPIb. Whether displayed as surface proteins on a phageid or
XX secreted as free scFv by Escherichia coli, the H1b scFv clones are
XX capable of inhibiting von Willebrand factor-dependent aggregation of
XX platelets. The scFv are composed of native human protein sequences and
XX are therefore attractive potential reagents for therapeutic purposes.
XX They provide a new class of antithrombotic agents, useful for the

Aap50091 Truncated
Aab07203 Human Fac
Aag38412 Arabidops
Aag38412 Arabidops
Aag29486 Arabidops
Aay95198 Anti-plat
Abp46004 Human Bly
Abp46027 Human Bly
Abp44926 Human Bly
Aao31136 Human CM0
Abg78329 Human Fv
Abg92026 Antibody
Abp30235 Streptoco
Abp45531 Human Bly
Abg78334 Human Fv
Abg92025 Antibody
Adc94375 E. faeciu
Aag32033 Arabidops
Adc97186 E. faeciu
Abg92020 Human ant
Aag29485 Arabidops

26 28 100.0 211 1 AAP50091
27 28 100.0 211 3 AAB07203
28 28 100.0 214 3 AAG38412
29 28 100.0 236 3 AAG29486
30 28 100.0 238 3 AAY95198
31 28 100.0 239 5 ABP46004
32 28 100.0 239 5 ABP46027
33 28 100.0 239 5 ABP44926
34 28 100.0 244 6 AAO31136
35 28 100.0 246 5 ABG78329
36 28 100.0 246 5 ABG92026
37 28 100.0 249 5 ABP30235
38 28 100.0 251 5 ABP45531
39 28 100.0 256 5 ABG78334
40 28 100.0 256 5 ABG92025
41 28 100.0 258 7 ADC94375
42 28 100.0 259 3 AAG32033
43 28 100.0 264 7 ADC97186
44 28 100.0 266 5 ABG92020
45 28 100.0 267 3 AAG29485

CC prevention of platelet-dependent thrombi in diseased arteries, bypass
 CC grafts, dialysis etc., and can also be used as diagnostic reagents;
 CC Methods of inhibiting aggregation of platelets, of binding human platelet
 CC GPIIb/IIIa and of selecting a VH or VL region of an antibody that
 CC inhibits platelet aggregation are claimed. Fragments of the scFv VH or VL
 CC chain, including CDR fragments, are also claimed
 XX
 XX Sequence 5 AA;

Query Match 100.0%; Score 28; DB 3; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 1 DYGMS 5

RESULT 2

ABG78239
 ID ABG78239 standard; peptide; 5 AA.

XX
 AC ABG78239;

XX
 DT 15-NOV-2002 (first entry)

XX Human Fv molecule hypervariable region related peptide #114.

XX Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
 KW disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
 KW lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.

XX Homo sapiens.

XX WO200259264-A2.

XX 01-AUG-2002.

XX 31-DEC-2001; 2001WO-US049440.

XX 29-DEC-2000; 2000US-00751181.

XX (BIOT-) BIO-TECHNOLOGY GEN CORP.

XX Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
 PI Plaksin D, Peretz T;
 XX WPI; 2002-619166/66.

XX Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
 PT or fragment, or construct of fragment with enhanced binding
 PT characteristics so as to selectively bind target cell in favor of other
 PT cells.

XX Claim 20; Page 208; 232pp; English.

XX The invention relates to a peptide or polypeptide comprising an Fv
 CC molecule, a construct or fragments or a construct of a fragment with
 CC enhanced binding characteristics which selectively and/or specifically
 CC binds to a target cell in favour of other cells, where binding is
 CC primarily determined by a first hypervariable region and Fv is a single
 CC chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
 CC association with or attached, coupled, combined, linked or fused to a
 CC pharmaceutical agent, is useful in the manufacture of a medicament, where
 CC the medicament has activity against a diseased cell, preferably a cancer
 CC cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
 CC myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
 CC acute myeloid leukaemia cell). The peptide is also useful for preparing a
 CC composition for use in inhibiting the growth of a diseased or cancer
 CC cell. This sequence represents a human Fv molecule hypervariable region
 CC related peptide of the invention

XX Sequence 5 AA;

Query Match 100.0%; Score 28; DB 5; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 1 DYGMS 5

RESULT 3

ABG91930
 ID ABG91930 standard; peptide; 5 AA.

XX
 AC ABG91930;

XX
 DT 04-DEC-2002 (first entry)

XX Human antibody fragment #114.

XX Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.

XX Homo sapiens.

XX WO200253700-A2.

XX 11-JUL-2002.

XX 31-DEC-2001; 2001WO-US049442.

XX 29-DEC-2000; 2000US-00751181.

XX 29-DEC-2000; 2000US-0258948P.

XX (BIOT-) BIO-TECHNOLOGY GEN CORP.

XX Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
 PI Szanton E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
 XX WPI; 2002-674776/72.

XX Novel isolated epitope present on cancer cells and important in
 PT physiological phenomena such as cell rolling, metastasis and
 PT inflammation, for treating autoimmune, inflammatory or cardiovascular
 PT diseases, and cancer.

XX Claim 25; Page 284; Opp; English.

XX The invention relates to an isolated epitope present on cancer cells and
 CC important in physiological phenomena such as cell rolling, metastasis and
 CC inflammation, where the epitope is capable of being bound by an antibody,
 CC its antigen-binding fragment or its complex comprising at least one
 CC antibody or its binding fragment having a first hypervariable region. The
 CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
 CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
 CC tumour or leukaemia cells, increase in number of tumour or leukaemia
 CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
 CC platelet and/or cell-platelet adhesion or aggregation, for increasing
 CC mortality of tumour or leukaemia cells, for increasing the susceptibility
 CC of diseased cells to damage by anti-disease, anti-cancer or anti-
 CC leukaemia agents, or for decreasing the number of tumour or leukaemia
 CC cells in a patient, or in the manufacture of a medicament for the above
 CC mentioned purposes. The epitopes are useful for diagnosing and treating
 CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
 CC diseases, cardiovascular diseases such as myocardial infarction,
 CC retinopathic diseases and other diseases mediated by abnormal platelet
 CC function and diseases caused by sulphated tyrosine-dependent protein-
 CC protein interactions. This sequence represents a human antibody fragment
 CC of the invention

SQ Sequence 5 AA;

Query Match 100.0%; Score 28; DB 5; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 DYGMS 5
 DB 1 DYGMS 5
 |||||

RESULT 4
 ID ABU11271 standard; peptide; 5 AA.
 XX AC ABU11271;
 XX DT 06-FEB-2003 (first entry)
 XX DE Human TANGO 268 VHCDR1 Peptide #5.
 XX KW Human; mouse; variable heavy; VH; antigen; cancer;
 KW complementarity determining region; TANGO 268; glycoprotein VI; GPVI;
 KW TANGO 268; extracellular matrix; collagen; platelet release;
 KW proliferation; migration; embryogenesis; inflammation; thrombosis;
 KW degranulation; thrombocytopaenia; antibody; thrombotic disorder;
 KW cerebral vascular disease; stroke; ischaemia; venous thromboembolism;
 KW leg swelling; pain; ulceration; pulmonary embolism; coronary disease;
 KW cardiovascular disease; angina pectoris; myocardial infarction;
 KW coronary restenosis; atherosclerosis; immunological disorder;
 KW developmental disorder; embryonic disorder; liver disorder;
 KW cerebral vascular disease; venous thromboembolism disease.
 XX OS Homo sapiens.
 XX WO200280968-A1.
 XX PD 17-OCT-2002.
 XX 09-APR-2002; 2002WO-US011122.
 XX 09-APR-2001; 2001US-00829495.
 XX (MILL-) MILLENNIUM PHARM INC.
 XX Busfield SJ, Valleval J, Jandrot-Ferrus M, Vainchenker W;
 PI Gill DS, Qian DW, Kingsbury G;
 PI WPI; 2003-058477/05.
 XX Novel substantially purified antibody immunospecifically binding to TANGO
 PT 268 antigen, useful for treating bleeding disorders such as
 PT thrombocytopenia, stroke, ischemia, pulmonary embolism, atherosclerosis.
 XX Claim 1; Page 111; 236pp; English.
 XX This invention relates to a novel purified antibody comprising a variable
 CC heavy (VH) complementarity determining region (CDR)1, VH CDR2 or VH CDR3;
 CC or variable light (VL) CDR1, VL CDR2 or VL CDR3, and immunospecifically
 CC binding to a TANGO 268 (also referred as glycoprotein VI (GPVI)) antigen.
 CC The antibodies of the invention act to decrease or block TANGO 268
 CC binding to extracellular matrix components, or as a collagen or platelet
 CC release and aggregation blocker. The antibodies of the invention are
 CC useful for modulating proliferation, migration, morphology,
 CC differentiation and/or function of megakaryocytes and platelets,
 CC including during development e.g. embryogenesis, modulating leukocyte-
 CC platelet and platelet-endothelium interactions in inflammation and/or
 CC thrombosis, and modulating platelet aggregation and degranulation. They
 CC are also useful for modulating disorders associated with abnormal or
 CC aberrant megakaryocyte and/or platelet proliferation, migration,
 CC morphology, differentiation and/or function, e.g. bleeding disorders such
 CC as thrombocytopaenia. Other diseases which may be modulated by these
 CC antibodies are thrombotic disorders, cerebral vascular diseases (e.g.

CC stroke and ischaemia) venous thromboembolism diseases (e.g. diseases
 CC involving leg swelling, pain and ulceration, pulmonary embolism, etc);
 CC coronary diseases (e.g. cardiovascular diseases including angina
 CC pectoris, myocardial infarction, coronary restenosis, atherosclerosis,
 CC etc); immunological disorders, developmental disorders, embryonic
 CC disorders, liver disorders, cerebral vascular diseases, venous
 CC thromboembolism disease, coronary diseases, and metastatic cancers. The
 CC antibodies of the invention only causes a transient decrease in platelet
 CC counts, platelet aggregation, and/or platelet activation and so have some
 CC advantages over prior art methods. The present sequence represents a
 CC peptide sequence used to generate the antibodies of the invention
 XX
 XX Sequence 5 AA;
 Query Match 100.0%; Score 28; DB 6; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 DYGMS 5
 DB 1 DYGMS 5
 |||||

RESULT 5
 ID AAR49460 standard; protein; 16 AA.
 XX AC AAR49460;
 XX DT 25-MAR-2003 (revised)
 DT 16-SEP-1994 (first entry)
 XX Factor VIII position 1175-1790.
 XX
 XX Naturally-occurring; immunomodulatory protein; human; therapy; class I;
 KW major histocompatibility complex; class II; allotype; type I diabetes;
 KW autoimmune disease; rheumatoid arthritis; T-cell-mediated response;
 KW multiple sclerosis; transplant rejection; vaccine; MHC.
 XX
 XX Homo sapiens.
 XX WO9404171-A1.
 XX PD 03-MAR-1994.
 XX 11-AUG-1993; 93WO-US007545.
 XX 11-AUG-1992; 92US-00925460.
 PR 15-JUN-1993; 93US-00077255.
 XX (HARD) HARVARD COLLEGE.
 PA Urban RG, Chicz RM, Vignali DA, Hedley ML, Stern LJ;
 PI Strominger JL;
 PI WPI; 1994-082825/10.
 XX Novel immunomodulatory peptide(s) and nucleic acids - useful for
 PT treatment of autoimmune diseases, transplant rejection and for
 PT vaccination.
 XX Disclosure; Page 41; 139pp; English.
 XX The sequences given in AAR49291-505 and AAR46981-7038 represent peptide
 CC fragments of naturally-occurring immunomodulatory proteins. These
 CC fragments are between 10-30 residues in length and bind to a human major
 CC histocompatibility complex (MHC) class II allotype. These peptides may be
 CC used for therapy of autoimmune diseases, such as type I diabetes,
 CC rheumatoid arthritis and multiple sclerosis, and to reduce transplant
 CC rejection. They may also be used for vaccination providing an exclusively
 CC T-cell-mediated response, which can be class I or class-II based, or
 CC both, depending on the length and character of the immunogenic peptides.
 CC (Updated on 25-MAR-2003 to correct PN field.) (Updated on 25-MAR-2003 to

CC correct PR field.)
 XX Sequence 16 AA;
 SQ

Query Match 100.0%; Score 28; DB 2; Length 16;
 Best Local Similarity 100.0%; Pred. No. 12;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 3 DYGMS 7

RESULT 6
 ABJ05002
 ID ABJ05002 standard; peptide; 20 AA.
 XX
 AC ABJ05002;
 XX
 DT 30-OCT-2002 (first entry)
 XX
 DE A3 peptide of fVII precursor SEQ ID No 58.
 KW Haemostatic; antibody inhibitor; factor VIII; T cell; immune response;
 KW haemophilia A; acquired haemophilia; human factor VIII.
 XX
 OS Homo sapiens.
 XX
 PN WO200260917-A2.
 XX
 PD 08-AUG-2002.
 XX
 PF 30-NOV-2001; 2001WO-US044945.
 XX
 PR 01-DEC-2000; 2000US-0250430P.
 XX
 PA (MINU) UNIV MINNESOTA.
 XX
 PI Conti-Fine BM;
 XX
 PS WPI; 2002-627462/67.
 XX
 SS The invention relates to isolated and purified peptides and variants thereof, as well as DNA encoding those peptides, useful for preventing or treating aberrant or pathogenic production of antibodies specific for factor VIII, particularly useful for treating hemophilia A or acquired hemophilia.
 XX
 PS Claim 44; Page 62; 120pp; English.
 XX
 SS The invention relates to isolated and purified peptides and variants thereof, as well as DNA encoding those peptides, useful to prevent or treat antibody inhibitors of factor VIII. The peptides are useful for preventing or inhibiting aberrant, pathogenic or undesirable antibody production or antibody binding that is specific for factor VIII. The peptides are also useful for preventing or inhibiting the priming or activity of T cells specific for factor VIII. These peptides are also useful for enhancing the activity or increasing the levels of modulatory T cells that inhibit the immune response to factor VIII. These peptides are useful in mammals, particularly in humans. The administration of these peptides does not increase the synthesis of a pathogenic antibody to factor VIII, or its biologically active fragment or functional equivalent. In particular, these peptides are useful for treating haemophilia A or acquired haemophilia. This sequence represents a human factor VIII peptide region relating to the invention

CC Sequence 20 AA;
 SQ

Query Match 100.0%; Score 28; DB 5; Length 20;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||

Db 9 DYGMS 13

RESULT 7
 AAP61435
 ID AAP61435 standard; protein; 26 AA.
 XX
 AC AAP61435;
 XX
 DT 25-MAR-2003 (revised)
 DT 02-AUG-1991 (first entry)
 XX
 DE Factor VIII:C cosuglant polypeptide.
 XX
 KW Factor VIII:C; coagulant; haemophilia; clotting; blood.
 XX
 OS Homo sapiens.
 XX
 PN EP182372-A.
 XX
 PD 28-MAY-1986.
 XX
 PF 19-NOV-1985; 85EP-00114743.
 XX
 PR 31-MAR-1983; 83US-00481105.
 PR 30-NOV-1983; 83US-00556508.
 PR 21-NOV-1984; 84US-00673916.
 PR 24-MAY-1985; 85US-00738134.
 PR 13-APR-1987; 87US-00037615.
 XX
 PA (SCRI) SCRIPPS CLINIC & RES FOUND.
 XX
 PI Zimmermann TS, Fulcher CA;
 XX
 DR WPI; 1986-138803/22.
 XX
 PT New factor VIII:c coagulant polypeptide(s) - useful for enhanced activity in treating haemophilia and obtd. by treating factor VIII:c with protease.
 PT
 XX
 PS Claim 6; Page 44; 46pp; English.
 XX
 SS The polypeptide has mol. wt. of approx. 79000-80000. It is an active Factor VIII:C coagulant polypeptide, and has enhanced coagulant activity. It is useful for treating haemophilia. It is also useful in the study and characterization of polypeptides or their complexes providing desired clotting behaviour in the blood of humans and other mammals. Activity may be 3-5 times that of known purified human factor VIII:C, or even 10-100 times. Activity is shown over a continuous period of at least 10 mins., and usually for much longer. The polypeptide may be mixed with other Factor VIII:C polypeptides to produce a specific coagulant activity of over 1000 units/ml. (Updated on 25-MAR-2003 to correct PA field.)

CC Sequence 26 AA;
 SQ

Query Match 100.0%; Score 28; DB 1; Length 26;
 Best Local Similarity 100.0%; Pred. No. 20;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 19 DYGMS 23

RESULT 8
 AAP50314
 ID AAP50314 standard; protein; 29 AA.
 XX
 AC AAP50314;
 XX
 DT 25-MAR-2003 (revised)
 DT 23-NOV-1991 (first entry)
 XX


```

DE Peptide encoded by porcine antihaemophilic factor exon (34-H1).
XX
XX Porcine antihaemophilic factor; haemophilia; factor VIIIC.
XX
OS Sus scrofa.
XX
XX WO8501961-A.
XX
XX PD 09-MAY-1985.
XX
XX PF 12-OCT-1984; 84WO-US001641.
XX
XX PR 28-OCT-1983; 83US-00546650.
XX
XX PR 24-AUG-1984; 84US-00644036.
XX
XX PA (GEMY ) GENETICS INST INC.
XX
XX PI Toole J;
XX
XX DR WPI; 1985-122479/20.
XX
XX PT Prodn. of human and porcine factor VIIIC - by using recombinant dna
XX techniques for their cellular prodn.
XX
XX PS Disclosure; Fig 1B; 71pp; English.
XX
XX CC The peptide is encoded by an exon of the porcine antihaemophilic factor
XX (AHF) gene. The exon, from nucleotides 169 to 267, encodes at least 30
XX amino acids from Phe 2 to Arg 31 of the 69 kD fragment. The AHF was obtd.
XX using anti-factor VIIIC monoclonal antibody. The peptide is useful for
XX the treatment of haemophilia. (Updated on 25-MAR-2003 to correct PR
XX field.) (Updated on 25-MAR-2003 to correct PA field.)
XX
XX SQ Sequence 29 AA;

Query Match 100.0%; Score 28; DB 1; Length 29;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 20 DYGMS 24
|||||

RESULT 9
AAP50318
ID AAP50318 standard; protein; 33 AA.
XX
XX AC AAP50318;
XX
XX DT 25-MAR-2003 (revised)
XX
XX DT 23-NOV-1991 (first entry)
XX
DE Peptide encoded by HaeIII insert 34-H1 bearing porcine antihaemophilic
DE factor exon.
XX
XX Porcine antihaemophilic factor; haemophilia; insert 34-H1; factor VIIIC.
XX
XX OS Sus scrofa.
XX
XX FH Key Location/Qualifiers
XX
XX FT Region 3..33
XX
XX FT /label= human factor VIIIC sequence
XX
XX PN WO8501961-A.
XX
XX PD 09-MAY-1985.
XX
XX PF 12-OCT-1984; 84WO-US001641.
XX
XX PR 28-OCT-1983; 83US-00546650.
XX
XX PR 24-AUG-1984; 84US-00644036.
XX
XX

DE Peptide encoded by porcine antihaemophilic factor exon (34-H1).
XX
XX Porcine antihaemophilic factor; haemophilia; factor VIIIC.
XX
OS Sus scrofa.
XX
XX WO8501961-A.
XX
XX PD 09-MAY-1985.
XX
XX PF 12-OCT-1984; 84WO-US001641.
XX
XX PR 28-OCT-1983; 83US-00546650.
XX
XX PR 24-AUG-1984; 84US-00644036.
XX
XX PA (GEMY ) GENETICS INST INC.
XX
XX PI Toole J;
XX
XX DR WPI; 1985-122479/20.
XX
XX PT Prodn. of human and porcine factor VIIIC - by using recombinant dna
XX techniques for their cellular prodn.
XX
XX PS Disclosure; Fig 1A; 71pp; English.
XX
XX CC The peptide is the N-terminal sequence of the 69 kD thrombin cleavage
XX prod. of porcine antihaemophilic factor (AHF). The AHF was obtd. using
XX anti-factor VIIIC monoclonal antibody. The peptide is useful for the
XX treatment of haemophilia. (Updated on 25-MAR-2003 to correct PR field.)
XX (Updated on 25-MAR-2003 to correct PA field.)
XX
XX SQ Sequence 33 AA;

Query Match 100.0%; Score 28; DB 1; Length 33;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 20 DYGMS 24
|||||

RESULT 10
AAP50313
ID AAP50313 standard; peptide; 41 AA.
XX
XX AC AAP50313;
XX
XX DT 25-MAR-2003 (revised)
XX
XX DT 23-NOV-1991 (first entry)
XX
DE Thrombin cleavage prod. of porcine antihaemophilic factor.
XX
XX Porcine antihaemophilic factor; haemophilia; thrombin cleavage prod;
XX factor VIIIC.
XX
XX OS Sus scrofa.
XX
XX PN WO8501961-A.
XX
XX PD 09-MAY-1985.
XX
XX PF 12-OCT-1984; 84WO-US001641.
XX
XX PR 28-OCT-1983; 83US-00546650.
XX
XX PR 24-AUG-1984; 84US-00644036.
XX
XX PA (GEMY ) GENETICS INST INC.
XX
XX PI Toole J;
XX
XX DR WPI; 1985-122479/20.
XX
XX PT Prodn. of human and porcine factor VIIIC - by using recombinant dna
XX techniques for their cellular prodn.
XX
XX PS Disclosure; Fig 1A; 71pp; English.
XX
XX CC The peptide is the N-terminal sequence of the 69 kD thrombin cleavage
XX prod. of porcine antihaemophilic factor (AHF). The AHF was obtd. using
XX anti-factor VIIIC monoclonal antibody. The peptide is useful for the
XX treatment of haemophilia. (Updated on 25-MAR-2003 to correct PR field.)
XX (Updated on 25-MAR-2003 to correct PA field.)
XX
XX SQ Sequence 33 AA;

Query Match 100.0%; Score 28; DB 1; Length 33;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 20 DYGMS 24
|||||

```

XX SQ Sequence 41 AA;

Query Match 100.0%; Score 28; DB 1; Length 41;
 Best Local Similarity 100.0%; Pred. No. 33;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 18 DYGMS 22

RESULT 11
 ABJ04948
 ID ABJ04948 standard; protein; 60 AA.
 XX
 AC ABJ04948;
 XX
 DT 30-OCT-2002 (first entry)
 XX
 DE A3 domain of fVII epitope SEQ ID No 4.
 XX
 KW Haemostatic; antibody inhibitor; factor VIII; T cell; immune response;
 KW haemophilia A; acquired haemophilia; human factor VIII.
 XX
 OS Homo sapiens.
 XX
 PN WC200260917-A2.
 XX
 PD 08-AUG-2002.
 XX
 PF 30-NOV-2001; 2001WO-US044945.
 XX
 PR 01-DEC-2000; 2000US-0250430P.
 XX
 PA (MINU) UNIV MINNESOTA.
 XX
 PI Conti-Pine BM;
 XX
 WPI; 2002-627462/67.
 XX
 New peptides or their variants, useful for preventing, treating or
 PT inhibiting aberrant or pathogenic production of antibodies specific for
 PT factor VIII, particularly useful for treating hemophilia A or acquired
 PT hemophilia.
 XX
 PS Claim 4; Page 84; 120pp; English.
 XX
 The invention relates to isolated and purified peptides and variants
 CC thereof, as well as DNA encoding those peptides, useful to prevent or
 CC treat antibody inhibitors of factor VIII. The peptides are useful for
 CC preventing or inhibiting aberrant, pathogenic or undesirable antibody
 CC production or antibody binding that is specific for factor VIII. The
 CC peptides are also useful for preventing or inhibiting the priming or
 CC activity of T cells specific for factor VIII. These peptides are also
 CC useful for enhancing the activity or increasing the levels of modulatory
 CC T cells that inhibit the immune response to factor VIII. These peptides
 CC are useful in mammals, particularly in humans. The administration of
 CC these peptides does not increase the synthesis of a pathogenic antibody
 CC to factor VIII, or its biologically active fragment or functional
 CC equivalent. In particular, these peptides are useful for treating
 CC haemophilia A or acquired haemophilia. This sequence represents a human
 CC factor VIII protein region relating to the invention

Sequence 60 AA;

Query Match 100.0%; Score 28; DB 5; Length 60;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 39 DYGMS 43

RESULT 12
 AAB07206
 ID AAB07206 standard; protein; 73 AA.
 XX
 AC AAB07206;
 XX
 DT 11-OCT-2000 (first entry)
 XX
 DE Human Factor VIIIC 77/80kd subunit partial protein sequence #1.
 XX
 KW Factor VIII C domain; human; blood clotting; haemophilia.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT Misc-difference 53
 FT /note= "encoded by CGT"
 XX
 PN EP1006182-A2.
 XX
 PD 07-JUN-2000.
 XX
 PF 11-JAN-1985; 2000EP-00200860.
 XX
 PR 12-JAN-1984; 84US-00570062.
 PR 26-OCT-1984; 84US-00664919.
 PR 11-JAN-1985; 85EP-00100223.
 PR 11-JAN-1985; 91EP-00113267.
 XX
 PA (CHIR) CHIRON CORP.
 PA (NOVO) NOVO-NORDISK AS.
 XX
 PI Kuo G, Rasmussen ME, Masiaz FR, Valenzuela P, Truett M;
 PI Favaloro J;
 XX
 WPI; 2000-367967/32.
 DR N-PSDB; AAB58444.
 XX
 DNA sequence of portion of human Factor VIIIC for treating and preventing
 PT the symptoms of hemophilia.
 XX
 Claim 3; Page 37; 39pp; English.
 XX
 The present sequence is the protein sequence for the 77/80kd subunit of
 CC human Factor VIIIC. Factor VIIIC is a plasma protein involved in blood
 CC coagulation, and is absent or defective in haemophilia A. The Factor
 CC VIIIC protein can, therefore, be used to treat haemophilia, as well as in
 CC the production of monoclonal antibodies to Factor VIIIC, and in
 CC diagnostic assays for the presence of Factor VIIIC subunits in
 CC physiological fluids, for example blood or serum

Sequence 73 AA;

Query Match 100.0%; Score 28; DB 3; Length 73;
 Best Local Similarity 100.0%; Pred. No. 62;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 60 DYGMS 64

RESULT 13
 AAB40073
 ID AAB40073 standard; protein; 98 AA.
 XX
 AC AAB40073;
 XX
 DT 05-FEB-2001 (first entry)
 XX
 DE Anti-hiL12 antibody H chain V region amino acid sequence SEQ ID 599.

XX Human; neutralising antibody; interleukin-12; IL-12; antiinflammatory;
 KW complementarity determining region; CDR; antirheumatic; antiarthritic;
 KW antisclerotic; neuroprotective; antipsoriatic; antiasthmatic; cardiant;
 KW antiparasitic; antibacterial; immunosuppressive; Crohn's disease;
 KW multiple sclerosis; rheumatoid arthritis.
 XX
 OS Homo sapiens.
 XX
 FN WO200056772-A1.
 XX
 XX 28-SEP-2000.
 PD
 XX 24-MAR-2000; 2000WO-US007946.
 PP
 XX 25-MAR-1999; 99US-0126603P.
 PR
 XX (BADI) BASF AG.
 PA (GEMY) GENETICS INST INC.
 XX
 XX Salfeld JG, Roguska M, Paskind M, Banerjee S, Tracey DE, White M;
 PI Kaymakalan Z, Labkovsky B, Sakorafas P, Friedrich S, Myles A;
 PI Verdan GW, Venturini A, Warne NW, Widom A, Elvin JG, Duncan AR;
 PI Derbyshire EG, Carmen S, Smith S, Holtet TL, Du Fou SL;
 XX
 DR WPI; 2000-638250/61.
 XX
 XX New human antibody specific for human interleukin-12 (IL-12) used to
 PT treat disorders characterized by aberrant IL-12 expression e.g. Crohn's
 PT disease and multiple sclerosis.
 XX
 PS Claim 75; Page 121; 377pp; English.
 XX
 CC This invention relates to a new human antibody specific for human
 CC interleukin-12 (IL-12). The invention also includes antigen binding
 CC portions that bind to IL-12. Sequences AAB39485-B39516 represent human
 CC anti-IL-12 antibody heavy and light chain complementarity determining
 CC region (CDR) amino acid sequences, and also includes variable region
 CC amino acid sequences. Other variable region amino acid sequences are
 CC given in AAB39517-B39560 and AAB40068-B40149. Sequences AAB39561-B39771
 CC represent anti-IL-12 CDR3 related amino acid sequences, AAB39772-B40063
 CC represent other CDR sequences. Light chain CDR3 consensus sequences are
 CC given in AAB40064-B40067. Primers used in the identification and
 CC construction of the antibodies of the invention are given in AAC61062-
 CC C61071. The antibody of the invention is a neutralising antibody and has
 CC antirheumatic; antiarthritic; antisclerotic; antiinflammatory;
 CC neuroprotective; antipsoriatic; antiasthmatic; cardiant; antiparasitic;
 CC antibacterial and immunosuppressive activity. The antibodies or antigen-
 CC binding fragments are useful in the treatment of disorders associated
 CC with detrimental release of human IL-12, especially Crohn's disease,
 CC multiple sclerosis and rheumatoid arthritis. They can also be used in the
 CC manufacture of a pharmaceutical composition to treat human IL-12
 CC disorders
 XX
 XX Sequence 98 AA;
 SQ
 Query Match 100.0%; Score 28; DB 3; Length 98;
 Best Local Similarity 100.0%; Pred. No. 86;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGMS 5
 DB 31 DYGMS 35
 RESULT 14
 ABG78186
 ID ABG78186 standard; protein; 98 AA.
 XX
 XX ABG78186;
 AC
 XX 15-NOV-2002 (first entry)
 DT
 XX

DE Human Fv molecule hypervariable region related peptide #61.
 XX
 XX Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
 KW disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
 KW lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.
 XX
 OS Homo sapiens.
 XX
 FN WO200259264-A2.
 XX
 XX 01-AUG-2002.
 PD
 XX 31-DEC-2001; 2001WO-US049440.
 PP
 XX 29-DEC-2000; 2000US-00751181.
 PR
 XX (BIOT-) BIO-TECHNOLOGY GEN CORP.
 PA
 XX Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
 PI Plaksin D, Peretz T;
 XX
 DR WPI; 2002-619166/66.
 XX
 PT Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
 PT or fragment, or construct of fragment with enhanced binding
 PT characteristics so as to selectively bind target cell in favor of other
 PT cells.
 XX
 PS Claim 13; Page 177-178; 232pp; English.
 XX
 CC The invention relates to a peptide or polypeptide comprising an Fv
 CC molecule, a construct or fragments or a construct of a fragment with
 CC enhanced binding characteristics which selectively and/or specifically
 CC binds to a target cell in favour of other cells, where binding is
 CC primarily determined by a first hypervariable region and Fv is a single
 CC chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
 CC association with or attached, coupled, combined, linked or fused to a
 CC pharmaceutical agent, is useful in the manufacture of a medicament, where
 CC the medicament has activity against a diseased cell, preferably a cancer
 CC cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
 CC myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
 CC acute myeloid leukaemia cell). The peptide is also useful for preparing a
 CC composition for use in inhibiting the growth of a diseased or cancer
 CC cell. This sequence represents a human Fv molecule hypervariable region
 CC related peptide of the invention
 XX
 SQ Sequence 98 AA;
 Query Match 100.0%; Score 28; DB 5; Length 98;
 Best Local Similarity 100.0%; Pred. No. 86;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGMS 5
 DB 31 DYGMS 35
 RESULT 15
 ABG91877
 ID ABG91877 standard; protein; 98 AA.
 XX
 XX ABG91877;
 AC
 XX 04-DEC-2002 (first entry)
 DT
 XX Human antibody fragment #61.
 DE
 XX Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.
 XX

OS Homo sapiens.
 XX WO200253700-A2.
 PN
 XX
 PD 11-JUL-2002.
 XX
 PF 31-DEC-2001; 2001WO-US049442.
 XX
 PR 29-DEC-2000; 2000US-00751181.
 PR 29-DEC-2000; 2000US-0258948P.
 XX
 XX (BIOT-) BIO-TECHNOLOGY GEN CORP.
 PA
 PA Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
 PI Szanthon E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
 XX WPI; 2002-674776/72.
 DR
 XX
 PT Novel isolated epitope present on cancer cells and important in
 PT physiological phenomena such as cell rolling, metastasis and
 PT inflammation, for treating autoimmune, inflammatory or cardiovascular
 PT diseases, and cancer.
 XX
 PS Disclosure; Page 255; Opp; English.
 XX
 CC The invention relates to an isolated epitope present on cancer cells and
 CC important in physiological phenomena such as cell rolling, metastasis and
 CC inflammation, where the epitope is capable of being bound by an antibody,
 CC its antigen-binding fragment or its complex comprising at least one
 CC antibody or its binding fragment having a first hypervariable region. The
 CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
 CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
 CC tumour or leukaemia cells, increase in number of tumour or leukaemia
 CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
 CC mortality of tumour or leukaemia cells, for increasing the susceptibility
 CC of diseased cells to damage by anti-disease, anti-cancer or anti-
 CC leukaemia agents, or for decreasing the number of tumour or leukaemia
 CC cells in a patient, or in the manufacture of a medicament for the above
 CC mentioned purposes. The epitopes are useful for diagnosing and treating
 CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
 CC diseases, cardiovascular diseases such as myocardial infarction,
 CC retinopathic diseases and other diseases mediated by abnormal platelet
 CC function and diseases caused by sulphated tyrosine-dependent protein-
 CC protein interactions. This sequence represents a human antibody fragment
 CC of the invention
 XX
 SQ Sequence 98 AA;
 Query Match 100.0%; Score 28; DB 5; Length 98;
 Best Local Similarity 100.0%; Pred. No. 86;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYQMS 5
 |||||
 Db 31 DYQMS 35

Search completed: September 18, 2004, 03:42:53
 Job time : 37.3571 secs

Query Match 100.0%; Score 28; DB 2; Length 16;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
|
|
|
|
Db 3 DYGMS 7

RESULT 2

US-08-488-379-170
; Sequence 170, Application US/08488379
; Patent No. 5880103

; GENERAL INFORMATION:
; APPLICANT: Robert G. Urban

; APPLICANT: Roman M. Chicz

; APPLICANT: Dario A. A. Vignali

; APPLICANT: Mary L. Hedley

; APPLICANT: Lawrence J. Stern

; APPLICANT: Jack L. Strominger

; TITLE OF INVENTION: IMMUNOMODULATORY PEPTIDES

; NUMBER OF SEQUENCES: 274

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson

; STREET: 225 Franklin Street

; CITY: Boston

; STATE: Massachusetts

; COUNTRY: U.S.A.

; ZIP: 02110-2804

; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb

; COMPUTER: IBM PS/2 Model 502 or 55SX

; OPERATING SYSTEM: MS-DOS (Version 5.0)

; SOFTWARE: WordPerfect (Version 5.1)

; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488,379

; FILING DATE:
; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/077,255

; FILING DATE: June 15, 1993

; APPLICATION NUMBER: 07/925,460

; FILING DATE: August 11, 1992

; ATTORNEY/AGENT INFORMATION:
; NAME: Clark, Paul T.

; REGISTRATION NUMBER: 30,162

; REFERENCE/DOCKET NUMBER: 00246/168001

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 542-5070

; TELEFAX: (617) 542-8906

; TELEX: 200154

; INFORMATION FOR SEQ ID NO: 170:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 16

; TYPE: amino acid

; STRANDEDNESS:
; TOPOLOGY: linear

US-08-488-379-170

Query Match 100.0%; Score 28; DB 2; Length 16;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
|
|
|
|
Db 3 DYGMS 7

RESULT 3

US-08-475-399A-170
; Sequence 170, Application US/08475399A

; Patent No. 6509031

; GENERAL INFORMATION:
; APPLICANT: Robert G. Urban

; APPLICANT: Roman M. Chicz

; APPLICANT: Dario A. A. Vignali

; APPLICANT: Mary L. Hedley

; APPLICANT: Lawrence J. Stern

; APPLICANT: Jack L. Strominger

; TITLE OF INVENTION: IMMUNOMODULATORY PEPTIDES

; NUMBER OF SEQUENCES: 273

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson

; STREET: 225 Franklin Street

; CITY: Boston

; STATE: Massachusetts

; COUNTRY: U.S.A.

; ZIP: 02110-2804

; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb

; APPLICANT: Urban, Robert G.
; APPLICANT: Chicz, Roman M.
; APPLICANT: Vignali, Dario A. A.
; APPLICANT: Hedley, Mary L.
; APPLICANT: Stern, Lawrence J.
; APPLICANT: Strominger, Jack L.
; TITLE OF INVENTION: IMMUNOMODULATORY PEPTIDES
; NUMBER OF SEQUENCES: 276
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson, P.C.

; STREET: 225 Franklin Street

; CITY: Boston

; STATE: MA

; COUNTRY: US

; ZIP: 02110-2804

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette

; COMPUTER: IBM Compatible

; OPERATING SYSTEM: Windows95

; SOFTWARE: FastSeq for Windows Version 2.0

; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/475,399A

; FILING DATE: 07-JUN-1995

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/077,255

; FILING DATE: 15-JUN-1993

; APPLICATION NUMBER: 07/925,460

; FILING DATE: 11-AUG-1992

; ATTORNEY/AGENT INFORMATION:
; NAME: Fraser, Janis K.

; REGISTRATION NUMBER: 34,819

; REFERENCE/DOCKET NUMBER: 00246/168003

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617/542-507

; TELEFAX: 617/542-890

; TELEX: 200154

; INFORMATION FOR SEQ ID NO: 170:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 16 amino acids

; TYPE: amino acid

; TOPOLOGY: linear

US-08-475-399A-170

Query Match 100.0%; Score 28; DB 4; Length 16;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
|
|
|
|
Db 3 DYGMS 7

RESULT 4

PCI-US93-07545-170

; Sequence 170, Application PC/TUS9307545
; GENERAL INFORMATION:
; APPLICANT: Robert G. Urban

; APPLICANT: Roman M. Chicz

; APPLICANT: Dario A. A. Vignali

; APPLICANT: Mary L. Hedley

; APPLICANT: Lawrence J. Stern

; APPLICANT: Jack L. Strominger

; TITLE OF INVENTION: IMMUNOMODULATORY PEPTIDES

; NUMBER OF SEQUENCES: 273

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson

; STREET: 225 Franklin Street

; CITY: Boston

; STATE: Massachusetts

; COUNTRY: U.S.A.

; ZIP: 02110-2804

; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb

COMPUTER: IBM PS/2 Model 50Z or 55SX
 OPERATING SYSTEM: MS-DOS (Version 5.0)
 SOFTWARE: WordPerfect (Version 5.1)
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: PCT/US93/07545
 FILING DATE: 19930811
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 07/925,460
 FILING DATE: August 11, 1992
 ATTORNEY/AGENT INFORMATION:
 NAME: Clark, Paul T.
 REGISTRATION NUMBER: 30,162
 REFERENCE/DOCKET NUMBER: 00246/168001
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (617) 542-5070
 TELEFAX: (617) 542-8906
 TELEX: 200154
 INFORMATION FOR SEQ ID NO: 170:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 16
 TYPE: amino acid
 STRANDEDNESS:
 TOPOLOGY: linear
 PCT-US93-07545-170

Query Match 100.0%; Score 28; DB 5; Length 16;
 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 3 DYGMS 7

RESULT 5
 US-08-652-816A-14
 Sequence 14, Application US/08652816A
 Patent No. 5872215

GENERAL INFORMATION:
 APPLICANT: Osbourn, JK
 APPLICANT: Allen, DJ
 APPLICANT: McCafferty, JG
 TITLE OF INVENTION: Specific binding members, materials and methods.
 TITLE OF INVENTION: methods.
 NUMBER OF SEQUENCES: 53
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
 STREET: 6300 Sears Tower, 233 South Wacker Drive
 CITY: Chicago
 STATE: Illinois
 COUNTRY: United States of America
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/652,816A
 FILING DATE: 23-MAY-1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: GB 9125579.4
 FILING DATE: 02-DEC-1991
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: GB 9125579.8
 FILING DATE: 02-DEC-1991
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: GB 9206318.9
 FILING DATE: 24-MAR-1992
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: GB 9206372.6
 FILING DATE: 23-SEP-1992
 PRIOR APPLICATION DATA:

APPLICATION NUMBER: GB 9525004.9
 FILING DATE: 07-DEC-1995
 PRIOR APPLICATION DATA: GB 9610824.6
 APPLICATION NUMBER: GB 9610824.6
 FILING DATE: 23-MAY-1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: PCT/GB92/02240
 FILING DATE: 02-DEC-1992
 PRIOR APPLICATION DATA: US 08/244,597
 FILING DATE: 01-JUN-1994
 ATTORNEY/AGENT INFORMATION:
 NAME: David W. Clough
 REGISTRATION NUMBER: 36,107
 REFERENCE/DOCKET NUMBER: 28111/33308
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 312-474-6300
 INFORMATION FOR SEQ ID NO: 14:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 116 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 US-08-652-816A-14

Query Match 100.0%; Score 28; DB 2; Length 116;
 Best Local Similarity 100.0%; Pred. No. 28;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 31 DYGMS 35

RESULT 6

US-08-545-809A-106
 Sequence 106, Application US/08545809A
 Patent No. 6096878
 GENERAL INFORMATION:
 APPLICANT: Honjo, Tasuko
 APPLICANT: Matsuda, Fumihiko
 TITLE OF INVENTION: HUMAN IMMUNOGLOBULIN VH GENE
 TITLE OF INVENTION: SEGMENTS AND DNA FRAGMENTS CONTAINING THE SAME
 NUMBER OF SEQUENCES: 145
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Fish & Richardson, P.C.
 STREET: 225 Franklin Street
 CITY: Boston
 STATE: MA
 COUNTRY: US
 ZIP: 02110-2804
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: Windows95
 SOFTWARE: FastSeq for Windows Version 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/545,809A
 FILING DATE: 27-MAR-1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: PCT/JP93/00603
 FILING DATE: 10-MAY-1993
 ATTORNEY/AGENT INFORMATION:
 NAME: Freeman, John W.
 REGISTRATION NUMBER: 29,066
 REFERENCE/DOCKET NUMBER: 06501/004001
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 617-542-5070
 TELEFAX: 617-542-8906
 TELEX: 200154
 INFORMATION FOR SEQ ID NO: 106:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 117 amino acids
 TYPE: amino acid

```
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-545-809A-106

Query Match      100.0%; Score 28; DB 3; Length 117;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
Db      50 DYGMS 54

RESULT 7
US-09-489-039A-8586
; Sequence 8586, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; PRIOR FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 8586
; LENGTH: 232
; TYPE: PRT
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-8586

Query Match      100.0%; Score 28; DB 4; Length 232;
Best Local Similarity 100.0%; Pred. No. 62;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
Db      219 DYGMS 223

RESULT 8
US-09-107-532A-4002
; Sequence 4002, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD/ROM ISO9660
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 6813:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 264 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHETICAL: YES
; ORIGINAL SOURCE:

; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-545-809A-106

Query Match      100.0%; Score 28; DB 3; Length 117;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
Db      50 DYGMS 54

RESULT 7
US-09-489-039A-8586
; Sequence 8586, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; PRIOR FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 8586
; LENGTH: 232
; TYPE: PRT
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-8586

Query Match      100.0%; Score 28; DB 4; Length 232;
Best Local Similarity 100.0%; Pred. No. 62;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
Db      219 DYGMS 223

RESULT 8
US-09-107-532A-4002
; Sequence 4002, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD/ROM ISO9660
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 6813:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 264 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHETICAL: YES
; ORIGINAL SOURCE:

; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-545-809A-106

Query Match      100.0%; Score 28; DB 3; Length 117;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
Db      50 DYGMS 54

RESULT 7
US-09-489-039A-8586
; Sequence 8586, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; PRIOR FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 8586
; LENGTH: 232
; TYPE: PRT
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-8586

Query Match      100.0%; Score 28; DB 4; Length 232;
Best Local Similarity 100.0%; Pred. No. 70;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
Db      172 DYGMS 176

RESULT 9
US-09-107-532A-6813
; Sequence 6813, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD/ROM ISO9660
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 6813:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 264 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHETICAL: YES
; ORIGINAL SOURCE:
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; ORGANISM: Enterococcus faecium
;
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (B) LOCATION 1...264
; SEQUENCE DESCRIPTION: SEQ ID NO: 6813
US-09-107-532A-6813

Query Match          100.0%; Score 28; DB 4; Length 264;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 178 DYGMS 182

RESULT 10
US-09-079-029-9
; Sequence 9, Application US/09079029
; Patent No. 6342369
; GENERAL INFORMATION:
; APPLICANT: Adams, Camilia W.
; APPLICANT: Ashkenazi, Avi J.
; APPLICANT: Chuntharapal, Anan
; APPLICANT: Kim, Kyung J.
; TITLE OF INVENTION: Apo-2 Receptor
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genentech, Inc.
; STREET: 1 DNA Way
; CITY: South San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94080
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: WinPatIn (Genentech)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/079,029
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Marschang, Diane L.
; REGISTRATION NUMBER: 35,600
; REFERENCE/DOCKET NUMBER: P1101R2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650/225-5416
; TELEFAX: 650/952-9881
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 309 amino acids
; TYPE: Amino Acid
; TOPOLOGY: Linear
US-09-079-029-9

Query Match          100.0%; Score 28; DB 4; Length 309;
Best Local Similarity 100.0%; Pred. No. 86;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 70 DYGMS 74

RESULT 11
US-08-724-984A-2
; Sequence 2, Application US/08724984A
; Patent No. 6388055
; GENERAL INFORMATION:
; APPLICANT: Derk Bergsma, Mary Brawner, and Usman Shabon
; TITLE OF INVENTION: NO. 6388055el Mouse Genomic Clone of the CC-
```

```
; TITLE OF INVENTION: CKR5 Receptor
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SmithKline Beecham Corporation
; STREET: 709 Swedeland Road, P.O. Box 1539
; CITY: King of Prussia
; STATE: PA
; COUNTRY: USA
; ZIP: 19406-0939
; COMPUTER READABLE FORM:
; MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE
; COMPUTER: IBM 486
; OPERATING SYSTEM: WINDOWS FOR WORKGROUPS
; SOFTWARE: MICROSOFT WORD
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/724,984A
; FILING DATE: October 3, 1996
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: William T. Han
; REGISTRATION NUMBER: 34,344
; REFERENCE/DOCKET NUMBER: ATG50023
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 610 270 5024
; TELEFAX: 610 270 5090
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 354
; TYPE: Amino Acid
; TOPOLOGY: Linear
US-08-724-984A-2

Query Match          100.0%; Score 28; DB 4; Length 354;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 15 DYGMS 19

RESULT 12
US-09-486-192-4
; Sequence 4, Application US/09486192
; Patent No. 6521440
; GENERAL INFORMATION:
; APPLICANT: Estell, David A.
; TITLE OF INVENTION: Proteases From Gram-Positive Organisms
; FILE REFERENCE: GC386-US
; CURRENT APPLICATION NUMBER: US/09/486,192
; CURRENT FILING DATE: 2000-02-22
; PRIOR APPLICATION NUMBER: PCT/US98/18677
; PRIOR FILING DATE: 1998-09-08
; PRIOR APPLICATION NUMBER: EP9719637.2
; PRIOR FILING DATE: 1997-09-15
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 4
; LENGTH: 474
; TYPE: PRT
; ORGANISM: Schizocaccharomyces pombe
US-09-486-192-4

Query Match          100.0%; Score 28; DB 4; Length 474;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 349 DYGMS 353
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RESULT 13
US-07-864-004B-6
; Sequence 6, Application US/07864004B
; Patent No. 5364771
; GENERAL INFORMATION:
; APPLICANT: Lollar, John S.
; APPLICANT: Runge, Marshall S.
; TITLE OF INVENTION: Hybrid Human/Porcine Factor VIII
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kilpatrick & Cody
; STREET: 1100 Peachtree Street
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: US
; ZIP: 30309
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/864,004B
; FILING DATE: 07 APRIL 1992
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284
; REFERENCE/DOCKET NUMBER: EMU106
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 404-815-6508
; TELEFAX: 404-815-6555
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 868 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: N-terminal
; ORIGINAL SOURCE:
; ORGANISM: Porcine
; US-07-864-004B-6
;
; Query Match 100.0%; Score 28; DB 1; Length 868;
; Best Local Similarity 100.0%; Pred. No. 2.8e+02;
; Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
; QY 1 DYGMS 5
; Db 804 DYGMS 808
;
; RESULT 14
US-08-251-937A-6
; Sequence 6, Application US/08251937A
; Patent No. 5583209
; GENERAL INFORMATION:
; APPLICANT: Lollar, John S.
; APPLICANT: Runge, Marshall S.
; TITLE OF INVENTION: Hybrid Human/Porcine Factor VIII
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kilpatrick & Cody
; STREET: 1100 Peachtree Street
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: US
; ZIP: 30309
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/251,937A
; FILING DATE: 07-APR-1992
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/864,004
; FILING DATE: 07-APR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284
; REFERENCE/DOCKET NUMBER: EMU106
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 404-815-6367
; TELEFAX: 404-815-6555
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 868 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: N-terminal
; ORIGINAL SOURCE:
; ORGANISM: Porcine
; US-08-251-937A-6
;
; Query Match 100.0%; Score 28; DB 1; Length 868;
; Best Local Similarity 100.0%; Pred. No. 2.8e+02;
; Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
; QY 1 DYGMS 5
; Db 804 DYGMS 808
;
; RESULT 15
US-08-212-133A-3
; Sequence 3, Application US/08212133A
; Patent No. 5683060
; GENERAL INFORMATION:
; APPLICANT: Lollar, John S.
; APPLICANT: Runge, Marshall S.
; TITLE OF INVENTION: Hybrid Human/Animal Factor VIII
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kilpatrick & Cody
; STREET: 100 Peachtree Street
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: US
; ZIP: 30303
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/212,133A
; FILING DATE: March 11, 1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/864,004
; FILING DATE: 07-APR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284
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Query Match 100.0%; Score 28; DB 1; Length 868;
Best Local Similarity 100.0%; Pred. No. 2.8e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 804 DYGMS 808

RESULT 15
US-08-212-133A-3
; Sequence 3, Application US/08212133A
; Patent No. 5683060
; GENERAL INFORMATION:
; APPLICANT: Lollar, John S.
; APPLICANT: Runge, Marshall S.
; TITLE OF INVENTION: Hybrid Human/Animal Factor VIII
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kilpatrick & Cody
; STREET: 100 Peachtree Street
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: US
; ZIP: 30303
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/212,133A
; FILING DATE: March 11, 1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/864,004
; FILING DATE: 07-APR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284

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/ REFERENCE/DOCKET NUMBER: EMU/76677
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 404-572-6508
/ TELEFAX: 404-572-6555
/ INFORMATION FOR SEQ ID NO: 3:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 868 amino acids
/ TYPE: amino acid
/ STRANDEDNESS: single
/ TOPOLOGY: linear
/ MOLECULE TYPE: protein
/ HYPOTHEICAL: YES
/ ANTI-SENSE: NO
/ FRAGMENT TYPE: N-terminal
/ ORIGINAL SOURCE:
/ ORGANISM: Porcine
/ FEATURE:
/ NAME/KEY: Protein
/ LOCATION: 1..868
/ OTHER INFORMATION: /note= "Predicted amino acid
/ OTHER INFORMATION: sequence of the B and part of the A2 domains of
/ OTHER INFORMATION: porcine factor VIII."
US-08-212-133A-3

Query Match 100.0%; Score 28; DB 1; Length 868;
Best Local Similarity 100.0%; Pred. No. 2.8e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 804 DYGMS 808
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Search completed: September 18, 2004, 03:48:46
Job time : 11.6071 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:46:39 ; Search time 198.571 Seconds
(without alignments)
8.086 Million cell updates/sec

Title: US-10-029-926B-114
Perfect score: 28
Sequence: 1 DYGMS 5

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1342398 seqs, 321133274 residues

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications AA:*

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- 2: /cgn2_6/ptodata/2/pubpaa/PCT_NEW PUB.pep.*
- 3: /cgn2_6/ptodata/2/pubpaa/US06_NEW PUB.pep.*
- 4: /cgn2_6/ptodata/2/pubpaa/US06_PUBCOMB.pep.*
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- 6: /cgn2_6/ptodata/2/pubpaa/PCTUS_PUBCOMB.pep.*
- 7: /cgn2_6/ptodata/2/pubpaa/US08_NEW PUB.pep.*
- 8: /cgn2_6/ptodata/2/pubpaa/US08_PUBCOMB.pep.*
- 9: /cgn2_6/ptodata/2/pubpaa/US09A_PUBCOMB.pep.*
- 10: /cgn2_6/ptodata/2/pubpaa/US09B_PUBCOMB.pep.*
- 11: /cgn2_6/ptodata/2/pubpaa/US09C_PUBCOMB.pep.*
- 12: /cgn2_6/ptodata/2/pubpaa/US09_NEW PUB.pep.*
- 13: /cgn2_6/ptodata/2/pubpaa/US10A_PUBCOMB.pep.*
- 14: /cgn2_6/ptodata/2/pubpaa/US10B_PUBCOMB.pep.*
- 15: /cgn2_6/ptodata/2/pubpaa/US10C_PUBCOMB.pep.*
- 16: /cgn2_6/ptodata/2/pubpaa/US10_NEW PUB.pep.*
- 17: /cgn2_6/ptodata/2/pubpaa/US60_NEW PUB.pep.*
- 18: /cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	5	9	US-09-832-312-73
2	28	100.0	5	11	US-09-829-495-73
3	28	100.0	5	12	US-10-029-926B-114
4	28	100.0	5	15	US-10-032-037B-114
5	28	100.0	5	15	US-10-029-988B-114
6	28	100.0	5	15	US-10-032-423A-114
7	28	100.0	20	16	US-10-433-273-58
8	28	100.0	60	16	US-10-433-273-4
9	28	100.0	98	12	US-10-453-698-60
10	28	100.0	98	12	US-10-029-926B-61
11	28	100.0	98	14	US-10-194-975-20
12	28	100.0	98	15	US-10-308-817-60
13	28	100.0	98	15	US-10-032-037B-61
14	28	100.0	98	15	US-10-029-988B-61
15	28	100.0	98	15	US-10-032-423A-61

16	28	100.0	98	16	US-10-379-392-21	Sequence 21, Appl
17	28	100.0	116	16	US-10-437-963-114282	Sequence 114282,
18	28	100.0	156	12	US-10-424-599-215325	Sequence 215325,
19	28	100.0	239	10	US-09-880-748-937	Sequence 937, App
20	28	100.0	239	10	US-09-880-748-2015	Sequence 2015, App
21	28	100.0	239	10	US-09-880-748-2038	Sequence 2038, App
22	28	100.0	239	12	US-10-293-418-937	Sequence 937, App
23	28	100.0	239	12	US-10-293-418-2015	Sequence 2015, App
24	28	100.0	239	12	US-10-293-418-2038	Sequence 2038, App
25	28	100.0	244	14	US-10-322-673-42	Sequence 42, Appl
26	28	100.0	251	10	US-09-880-748-1542	Sequence 1542, App
27	28	100.0	251	12	US-10-293-418-1542	Sequence 1542, App
28	28	100.0	266	15	US-10-032-037B-204	Sequence 204, App
29	28	100.0	266	15	US-10-029-988B-204	Sequence 204, App
30	28	100.0	266	15	US-10-032-423A-204	Sequence 204, App
31	28	100.0	276	12	US-10-425-114-70003	Sequence 70003, A
32	28	100.0	277	12	US-10-029-926B-25	Sequence 25, Appl
33	28	100.0	277	12	US-10-029-926B-203	Sequence 203, Appl
34	28	100.0	277	15	US-10-032-037B-25	Sequence 25, Appl
35	28	100.0	277	15	US-10-032-037B-203	Sequence 203, App
36	28	100.0	277	15	US-10-029-988B-25	Sequence 25, Appl
37	28	100.0	277	15	US-10-029-988B-203	Sequence 203, App
38	28	100.0	277	15	US-10-032-423A-25	Sequence 25, Appl
39	28	100.0	279	12	US-10-032-423A-203	Sequence 203, App
40	28	100.0	279	12	US-10-425-114-36951	Sequence 36951, A
41	28	100.0	299	16	US-10-767-701-44522	Sequence 44522, A
42	28	100.0	300	12	US-10-425-114-37904	Sequence 37904, A
43	28	100.0	309	13	US-10-052-798-9	Sequence 9, Appl
44	28	100.0	309	14	US-10-288-917-9	Sequence 9, Appl
45	28	100.0	309	15	US-10-423-448-9	Sequence 9, Appl

ALIGNMENTS

RESULT 1
US-09-832-312-73
; Sequence 73, Application US/09832312
; Patent No. US20010049829A1
; GENERAL INFORMATION:
; APPLICANT: Busfield et al.
; TITLE OF INVENTION: GLYCOPROTEIN VI AND USES THEREOF
; FILE REFERENCE: 7853-234
; CURRENT APPLICATION NUMBER: US/09/832,312
; PRIOR FILING DATE: 2001-04-09
; PRIOR APPLICATION NUMBER: 09/610,118
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: 09/503,387
; PRIOR FILING DATE: 2000-02-14
; PRIOR APPLICATION NUMBER: 09/454,824
; PRIOR FILING DATE: 1999-12-06
; PRIOR APPLICATION NUMBER: 09/345,468
; PRIOR FILING DATE: 1999-06-30
; NUMBER OF SEQ ID NOS: 78
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 73
; LENGTH: 5
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-832-312-73

Query Match 100.0%; Score:28; DB 9; Length 5;
Best Local Similarity 100.0%; Pred.No. 1.2e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 1 DYGMS 5

RESULT 2
US-09-829-495-73
; Sequence 73, Application US/09829495

us-10-029-926b-114.rapb

Mon Sep 20 10:58:51 2004

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; Publication No. US20040001826A1
; GENERAL INFORMATION:
; APPLICANT: Busfield SJ
; APPLICANT: Villevall J
; APPLICANT: Jandrot-Perrus M
; APPLICANT: Vainchenker W
; APPLICANT: Gill DS
; APPLICANT: Qian MD
; TITLE OF INVENTION: GLYCOPROTEIN VI AND USES THEREOF
; FILE REFERENCE: 7853-234
; CURRENT APPLICATION NUMBER: US/09/829,495
; CURRENT FILING DATE: 2001-04-09
; PRIOR APPLICATION NUMBER: 09/610,118
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: 09/503,387
; PRIOR FILING DATE: 2000-02-14
; PRIOR APPLICATION NUMBER: 09/454,824
; PRIOR FILING DATE: 1999-12-06
; PRIOR APPLICATION NUMBER: 09/345,468
; PRIOR FILING DATE: 1999-06-30
; NUMBER OF SEQ ID NOS: 78
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 73
; LENGTH: 5
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-829-495-73

Query Match      100.0%; Score 28; DB 11; Length 5;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      1 DYGMS 5

RESULT 3
US-10-029-926B-114
; Sequence 114, Application US/10029926B
; Publication No. US20040073011A1
; GENERAL INFORMATION:
; APPLICANT: HAGAY, et al.
; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
; FILE REFERENCE: 10793/50
; CURRENT APPLICATION NUMBER: US/10/029,926B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 203
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 114
; LENGTH: 5
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-029-926B-114

Query Match      100.0%; Score 28; DB 12; Length 5;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      1 DYGMS 5

RESULT 4
US-10-032-037B-114
; Sequence 114, Application US/10032037B
; Publication No. US20040001822A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/45
; CURRENT APPLICATION NUMBER: US/10/032,423A
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 114
; LENGTH: 5
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-032-423A-114

Query Match      100.0%; Score 28; DB 15; Length 5;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      1 DYGMS 5

RESULT 5
US-10-029-988B-114
; Sequence 114, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 114
; LENGTH: 5
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-029-988B-114

Query Match      100.0%; Score 28; DB 15; Length 5;
Best Local Similarity 100.0%; Pred. No. 1.2e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      1 DYGMS 5

RESULT 6
US-10-032-423A-114
; Sequence 114, Application US/10032423A
; Publication No. US20040002450A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/45
; CURRENT APPLICATION NUMBER: US/10/032,423A
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 114
; LENGTH: 5
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-032-423A-114

```

Query Match 100.0%; Score 28; DB 15; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.2e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 1 DYGMS 5

RESULT 7

US-10-433-273-58
 ; Sequence 58, Application US/10433273
 ; Publication No. US20040096456A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Regents of the University of Minnesota
 ; APPLICANT: Conti-Fine, Bianca M.
 ; TITLE OF INVENTION: Methods to Treat Hemophilia
 ; FILE REFERENCE: 600.507W01
 ; CURRENT APPLICATION NUMBER: US/10/433,273
 ; CURRENT FILING DATE: 2003-11-17
 ; PRIOR APPLICATION NUMBER: US 60/250,430
 ; PRIOR FILING DATE: 2000-12-01
 ; NUMBER OF SEQ ID NOS: 61
 ; SEQ ID NO 58
 ; LENGTH: 20
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-10-433-273-58

Query Match 100.0%; Score 28; DB 16; Length 20;
 Best Local Similarity 100.0%; Pred. No. 25;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 9 DYGMS 13

RESULT 8

US-10-433-273-4
 ; Sequence 4, Application US/10433273
 ; Publication No. US20040096456A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Regents of the University of Minnesota
 ; APPLICANT: Conti-Fine, Bianca M.
 ; TITLE OF INVENTION: Methods to Treat Hemophilia
 ; FILE REFERENCE: 600.507W01
 ; CURRENT APPLICATION NUMBER: US/10/433,273
 ; CURRENT FILING DATE: 2003-11-17
 ; PRIOR APPLICATION NUMBER: US 60/250,430
 ; PRIOR FILING DATE: 2000-12-01
 ; NUMBER OF SEQ ID NOS: 61
 ; SEQ ID NO 4
 ; LENGTH: 60
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-10-433-273-4

Query Match 100.0%; Score 28; DB 16; Length 60;
 Best Local Similarity 100.0%; Pred. No. 77;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 39 DYGMS 43

RESULT 9

US-10-453-698-60
 ; Sequence 60, Application US/10453698
 ; Publication No. US20040038308A1
 ; GENERAL INFORMATION:

; APPLICANT: Rother, Russell
 ; TITLE OF INVENTION: HYBRID ANTIBODIES
 ; FILE REFERENCE: 82 CIP (1087-37 CIP)
 ; CURRENT APPLICATION NUMBER: US/10/453,698
 ; CURRENT FILING DATE: 2003-06-03
 ; NUMBER OF SEQ ID NOS: 196
 ; SOFTWARE: Patent in version 3.2
 ; SEQ ID NO 60
 ; LENGTH: 98
 ; TYPE: PRT
 ; ORGANISM: human
 US-10-453-698-60

Query Match 100.0%; Score 28; DB 12; Length 98;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 31 DYGMS 35

RESULT 10

US-10-029-926B-61
 ; Sequence 61, Application US/10029926B
 ; Publication No. US20040073011A1
 ; GENERAL INFORMATION:
 ; APPLICANT: HAGAY, et al.
 ; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
 ; FILE REFERENCE: 10793/50
 ; CURRENT APPLICATION NUMBER: US/10/029,926B
 ; CURRENT FILING DATE: 2001-12-31
 ; PRIOR APPLICATION NUMBER: 60/258,948
 ; PRIOR FILING DATE: 12/29/2000
 ; NUMBER OF SEQ ID NOS: 203
 ; SOFTWARE: FastSeq for Windows Version 3.0
 ; SEQ ID NO 61
 ; LENGTH: 98
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-10-029-926B-61

Query Match 100.0%; Score 28; DB 12; Length 98;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 31 DYGMS 35

RESULT 11

US-10-194-975-20
 ; Sequence 20, Application US/10194975
 ; Publication No. US20030039649A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Foote, Jefferson
 ; TITLE OF INVENTION: Super Humanized Antibodies
 ; FILE REFERENCE: 501231.01
 ; CURRENT APPLICATION NUMBER: US/10/194,975
 ; CURRENT FILING DATE: 2002-10-10
 ; PRIOR APPLICATION NUMBER: US 60/305,111
 ; PRIOR FILING DATE: 2001-07-12
 ; NUMBER OF SEQ ID NOS: 122
 ; SOFTWARE: Patent in version 3.1
 ; SEQ ID NO 20
 ; LENGTH: 98
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-10-194-975-20

Query Match 100.0%; Score 28; DB 14; Length 98;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;

us-10-029-926b-114.rapb

Mon Sep 20 10:58:51 2004

```

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 31 DYGMS 35

RESULT 12
US-10-308-817-60
; Sequence 60, Application US/10308817
; Publication No. US20030219861A1
; GENERAL INFORMATION:
; APPLICANT: Rother, Russell
; TITLE OF INVENTION: HYBRID ANTIBODIES
; FILE REFERENCE: 1087-37
; CURRENT APPLICATION NUMBER: US/10/308,817
; CURRENT FILING DATE: 2002-12-03
; NUMBER OF SEQ ID NOS: 195
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 60
; LENGTH: 98
; TYPE: PRT
; ORGANISM: human
US-10-308-817-60

Query Match 100.0%; Score 28; DB 15; Length 98;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 31 DYGMS 35

RESULT 13
US-10-032-037B-61
; Sequence 61, Application US/10032037B
; Publication No. US20040001822A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/44
; CURRENT APPLICATION NUMBER: US/10/032,037B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 61
; LENGTH: 98
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-61

Query Match 100.0%; Score 28; DB 15; Length 98;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 31 DYGMS 35

RESULT 14
US-10-029-988B-61
; Sequence 61, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 61
; LENGTH: 98
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-61

Query Match 100.0%; Score 28; DB 15; Length 98;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 31 DYGMS 35

Search completed: September 18, 2004, 04:20:42
Job time : 199.571 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:33:04 ; Search time 10,1786 Seconds
(without alignments)
47,252 Million cell updates/sec

Title: US-10-029-926b-114

Perfect score: 28

Sequence: 1 DYGMS 5

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: Pir1.*

2: Pir2.*

3: Pir3.*

4: Pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	98	2 S26928	Ig heavy chain V r
2	28	100.0	112	2 PH1654	Ig heavy chain V r
3	28	100.0	119	2 A43413	Ig heavy chain V r
4	28	100.0	227	2 AB1076	probable fibrial
5	28	100.0	268	2 H84684	En/Spm-like transp
6	28	100.0	268	2 T04660	hypothetical prote
7	28	100.0	334	2 H95307	arginine deiminase
8	28	100.0	340	2 H72681	hypothetical prote
9	28	100.0	411	2 S74760	hypothetical prote
10	28	100.0	448	2 AB3043	oxidoreductase Atu
11	28	100.0	468	2 B98243	hypothetical prote
12	28	100.0	474	2 T38737	hypothetical prote
13	28	100.0	478	2 F89651	probable CMAX pren
14	28	100.0	502	2 T24471	protein T04F8.2 [i
15	28	100.0	609	2 H71285	hypothetical prote
16	28	100.0	650	2 G87572	probable cell divi
17	28	100.0	756	2 D96527	calcium-binding pr
18	28	100.0	830	2 I50455	protein F27J15.24
19	28	100.0	869	2 A25945	prolactin receptor
20	28	100.0	916	2 B84473	coagulation factor
21	28	100.0	1008	2 H85055	copla-like retroel
22	28	100.0	1250	2 T27706	probable transpos
23	28	100.0	2133	2 T42763	hypothetical prote
24	28	100.0	2319	2 A47004	coagulation factor
25	28	100.0	2351	1 B2HU	coagulation factor
26	25	89.3	109	2 T47696	hypothetical prote
27	25	89.3	113	2 B36259	Ig heavy chain V r
28	25	89.3	136	2 S35759	BHD9D10 protein -
29	25	89.3	138	1 D69979	conserved hypothet

30	25	89.3	145	2 D69383	conserved hypothet
31	25	89.3	154	2 B61027	hypothetical prote
32	25	89.3	171	2 S69895	helicase (BC 3.6.1
33	25	89.3	188	2 T48671	extracellular heme
34	25	89.3	203	2 C85288	hypothetical prote
35	25	89.3	203	2 T05519	hypothetical prote
36	25	89.3	220	2 H81048	biopolymer transpo
37	25	89.3	233	1 F63178	conserved hypothet
38	25	89.3	238	2 H70734	hypothetical prote
39	25	89.3	250	2 C85040	hypothetical prote
40	25	89.3	251	2 D69861	RNA polymerase sig
41	25	89.3	264	2 AF3550	phosphoglycolate p
42	25	89.3	277	2 S76396	hypothetical prote
43	25	89.3	289	2 A13166	hypothetical prote
44	25	89.3	291	2 A63545	mRNA 3'-end proces
45	25	89.3	298	2 T29685	hypothetical prote

ALIGNMENTS

RESULT 1

S26928 Ig heavy chain V region (DP-32) - human (fragment)

C;Species: Homo sapiens (man)

C;Date: 22-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 23-Jul-1999

C;Accession: S26928

R;Tomlinson, I.M.; Walter, G.; Marks, J.D.; Llewelyn, M.B.; Winter, G.

J. Mol. Biol. 227, 776-798, 1992

A;Title: The repertoire of human germline V(H) sequences reveals about fifty groups of

A;Reference number: S26985; MUID:93021117; PMID:1404388

A;Accession: S26928

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-98 <TCM>

A;Cross-references: EMBL:Z12334; NID:932887; PIDN:CAA78204.1; PID:932888

C;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotetramer; immunoglobulin

F;15-98/Domain: immunoglobulin homology <IMV>

Query Match 100.0%; Score 28; DB 2; Length 98;

Best Local Similarity 100.0%; Pred.No.18;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5

Db 31 DYGMS 35

RESULT 2

PH1654

Ig heavy chain V region (clone 6H9) - human (fragment)

C;Species: Homo sapiens (man)

C;Date: 24-Feb-1994 #sequence_revision 24-Feb-1994 #text_change 16-Aug-1996

C;Accession: PH1654

R;Hillson, J.L.; Karr, N.S.; Opplinger, I.R.; Mannik, M.; Sasso, E.H.

J. Exp. Med. 178, 331-336, 1993

A;Title: The structural basis of germline-encoded VH3 immunoglobulin binding to staphyl

A;Reference number: PH1642; MUID:93301610; PMID:8315388

A;Accession: PH1654

A;Molecule type: mRNA

A;Residues: 1-112 <HLL>

A;Experimental source: B cell

C;Superfamily: immunoglobulin V region; immunoglobulin homology

C;Keywords: heterotetramer; immunoglobulin

F;7-90/Domain: immunoglobulin homology <IMV>

Query Match 100.0%; Score 28; DB 2; Length 112;

Best Local Similarity 100.0%; Pred.No.21;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5

|||||

Db 23 DYGMs 27

RESULT 3

A43413

Ig heavy chain V region - mouse (fragment)

A:Accession: H84684

C:Species: Mus musculus (house mouse)

C>Date: 27-Apr-1993 #sequence_revision 18-Nov-1994 #text_change 16-Aug-1996

C:Accession: A43413

R:Tomiyama, Y.; Brojer, E.; Ruggeri, Z.M.; Shattil, S.J.; Smiltneck, J.; Gorski, J.; Kum

J. Biol. Chem. 267, 18085-18092, 1992

A:Title: A molecular model of RGD ligands. Antibody D gene segments that direct specific

A:Reference number: A43413; MUID:92388177; PMID:1517241

A:Accession: A43413

A>Status: preliminary; not compared with conceptual translation

A:Molecule type: nucleic acid

A:Residues: 1-119 <TOM>

A>Note: sequence extracted from NCBI backbone (NCBIP:112815)

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

P:9-92/Domain: immunoglobulin homology <IMM>

Query Match 100.0%; Score 28; DB 2; Length 119;

Best Local Similarity 100.0%; Pred. No. 22; Mismatches 0; Indels 0; Gaps 0;

Matches 5; Conservative 0;

QY 1 DYGMs 5

Db 25 DYGMs 29

RESULT 4

AB1076

probable fibrillar chaperone protein sthB [imported] - Salmonella enterica subsp. enteric

C:Species: Salmonella enterica subsp. enterica serovar Typhi

A>Note: this species has also been called Salmonella typhi

C>Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 18-Nov-2002

C:Accession: AB1076

R:Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; Wain, J.; Churcher,

th, T.; Conerton, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.; White, N.; Farrar,

S.; Moule, S.; O'Gaora, P.

Nature 413, 848-852, 2001

A:Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.

A:Title: Complete genome sequence of a multiple drug resistant salmonella enterica serov

A:Reference number: AB0502; MUID:21534947; PMID:11677608

A:Accession: AB1076

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-227 <PAR>

A:Cross-references: GB:AU513382; PIDN:CAD03425.1; PID:gl6505691; GSPDB:GN00176

C:Genetics:

A:Gene: sthB

C:Superfamily: chaperone protein papD

Query Match 100.0%; Score 28; DB 2; Length 227;

Best Local Similarity 100.0%; Pred. No. 45;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMs 5

Db 215 DYGMs 219

RESULT 5

H84684

En/Spm-like transposon protein [imported] - Arabidopsis thaliana

C:Species: Arabidopsis thaliana (mouse-ear cress)

C>Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001

C:Accession: H84684

R:Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;

M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; Vanaken, S.E.; Umayam, L.; Tallon, L.

euss, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, J

Nature 402, 761-768, 1999

A:Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.

A:Reference number: A84420; MUID:20083487; PMID:10617197

A:Accession: H84684

A>Status: Preliminary

A:Molecule type: DNA

A:Residues: 1-268 <STO>

A:Cross-references: GB:AB002093; NID:94432833; PIDN:AD20682.1; GSPDB:GN00139

C:Genetics:

A:Gene: At2G28440

A:Map position: 2

Query Match 100.0%; Score 28; DB 2; Length 268;

Best Local Similarity 100.0%; Pred. No. 54;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMs 5

Db 218 DYGMs 222

RESULT 6

T04660

hypothetical protein F8D20.60 - Arabidopsis thaliana

C:Species: Arabidopsis thaliana (mouse-ear cress)

C>Date: 23-Apr-1999 #sequence_revision 23-Apr-1999 #text_change 21-May-1999

C:Accession: T04660

R:Bevan, M.; Rose, M.; Hempel, S.; Entian, K.D.; Jesse, T.; Heijnen, L.; Vos, P.; Mewes,

submitted to the Protein Sequence Database, July 1998

A:Reference number: Z15381

A:Accession: T04660

A:Molecule type: DNA

A:Residues: 1-268 <BEV>

A:Cross-references: EMBL:AL011135

A:Experimental source: cultivar Columbia; BAC clone F8D20

C:Genetics:

A:Map position: 4

A:Introns: 75/1; 241/1

A>Note: F8D20.60

Query Match 100.0%; Score 28; DB 2; Length 268;

Best Local Similarity 100.0%; Pred. No. 54;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMs 5

Db 263 DYGMs 267

RESULT 7

H93307

arginine deiminase (EC 3.5.3.6) ArcB [imported] - Sinorhizobium meliloti (strain 1021)

C:Species: Sinorhizobium meliloti

C>Date: 24-Aug-2001 #sequence_revision 24-Aug-2001 #text_change 14-Sep-2001

C:Accession: H93307

R:Barnett, M.J.; Fisher, R.F.; Jones, T.; Komp, C.; Abola, A.P.; Barloy-Hubler, F.; Rowe

; Kallman, S.; Keating, D.H.; Palm, C.; Peck, M.C.; Surzycki, R.; Wells, D.H.; Yeh, K.C.

Proc. Natl. Acad. Sci. U.S.A. 98, 9883-9888, 2001

A:Title: Nucleotide sequence and predicted functions of the entire Sinorhizobium melilot

A:Reference number: A95262; MUID:21396509; PMID:11481432

A:Accession: H93307

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-334 <KUR>

A:Cross-references: GB:AE006469; PIDN:AAK5026.1; PID:gl4523456; GSPDB:GN00165

A:Experimental source: strain 1021, megaplasmid pSymA

R:Galibert, F.; Finan, T.M.; Long, S.R.; Puhler, A.; Abola, P.; Ampe, F.; Barloy-Hubler,

Pel, D.; Chain, P.; Cowie, A.; Davis, R.W.; Dreano, S.; Federspiel, N.A.; Fisher, R.F.;

L.; Hyman, R.W.; Jones, T.

Science 283, 668-672, 2001

A:Authors: Kahn, D.; Kahn, M.L.; Kallman, S.; Keating, D.H.; Kiss, E.; Komp, C.; Leilaure,

hebaull, P.; Vandenbol, M.; Vorholter, F.J.; Weidner, S.; Wells, D.H.; Wong, K.; Yeh, K.

A:Title: The composite genome of the legume symbiont Sinorhizobium meliloti.

A:Reference number: A96039; MUID:21368234; PMID:11474104

```

A;Contents: annotation
C;Genetics:
A;Gene: arcB
A;Genome: plasmid
C;Superfamily: ornithine carbamoyltransferase; aspartate/ornithine carbamoyltransferase
C;Keywords: hydrolase

Query Match      100.0%; Score 28; DB 2; Length 334;
Best Local Similarity 100.0%; Pred. No. 68;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      292 DYGMS 296

RESULT 8
H72681
hypothetical protein APE0874 - Aeropyrum pernix (strain K1)
C;Species: Aeropyrum pernix
C;Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Jun-2000
C;Accession: H72681
R;Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takahawa, H.; Takamiya, M.; Masuda, S.; Furuhashi, T.; Tanaka, T.; Kudo, Y.; Yamazaki, J.; KDNA Res. 6, 83-101, 1999
A;Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyrum pernix
A;Reference number: A72450; MUID:99310339; PMID:10382966
A;Accession: H72681
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-340 <KAW>
A;Cross-references: DDBJ:AP000060; NID:G5104188; PIDN:BAA79856.1; PID:G5104541
A;Experimental source: strain K1
C;Genetics:
A;Gene: APE0874
C;Superfamily: Aeropyrum pernix hypothetical protein APE0874

Query Match      100.0%; Score 28; DB 2; Length 340;
Best Local Similarity 100.0%; Pred. No. 69;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      82 DYGMS 86

RESULT 9
S74760
hypothetical protein slr1617 - Synechocystis sp. (strain PCC 6803)
C;Species: Synechocystis sp.
A;Variety: PCC 6803
C;Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 08-Oct-1999
C;Accession: S74760
R;Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima, N.; O.K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yasuda DNA Res. 3, 109-136, 1996
A;Title: Sequence analysis of the genome of the unicellular cyanobacterium Synechocystis sp.
A;Reference number: S74322; MUID:97061201; PMID:8905231
A;Accession: S74760
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-411 <KAN>
A;Cross-references: EMBL:D90901; GB:A8001339; NID:G1651897; PIDN:BAA16911.1; PID:dl01764
A;Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996

Query Match      100.0%; Score 28; DB 2; Length 411;
Best Local Similarity 100.0%; Pred. No. 86;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      65 DYGMS 69

```

```

RESULT 10
AB3043
oxidoreductase Atu3958 [imported] - Agrobacterium tumefaciens (strain C58, Dupont)
C;Species: Agrobacterium tumefaciens
C;Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 18-Nov-2002
C;Accession: AB3043
R;Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo, J.; Erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutyavin, T.; Levy, R.; Li, M.; McClellan, K.; Karp, P.; Romero, P.; Zhang, S.
Science 294, 2317-2323, 2001
A;Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm, ster, E.W.
A;Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
A;Reference number: AB2577; MUID:21608550; PMID:11743193
A;Accession: AB3043
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-448 <KUR>
A;Cross-references: PIDN:AAL44760.1; PID:G17742396; GSPDB:GN00187
A;Experimental source: strain C58 (Dupont)
C;Genetics:
A;Gene: Atu3958
A;Map position: linear chromosome

Query Match      100.0%; Score 28; DB 2; Length 448;
Best Local Similarity 100.0%; Pred. No. 94;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      124 DYGMS 128

RESULT 11
B98243
hypothetical protein AGR_L1791 [imported] - Agrobacterium tumefaciens (strain C58, Cer
C;Species: Agrobacterium tumefaciens
C;Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 18-Nov-2002
C;Accession: B98243
R;Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Quorillo, B.; Goldman A.; Liu, F.; Wollan, C.; Allinger, M.; Doughty, D.; Scott, C.; Lappas, C.; Markelz, B. Science 294, 2323-2328, 2001
A;Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tu
A;Reference number: A97359; MUID:21608551; PMID:11743194
A;Accession: B98243
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-468 <KUR>
A;Cross-references: GB:AE007870; PIDN:AAK9468.1; PID:G15159336; GSPDB:GN00170
C;Genetics:
A;Gene: AGR_L1791
A;Map position: linear chromosome

Query Match      100.0%; Score 28; DB 2; Length 468;
Best Local Similarity 100.0%; Pred. No. 99;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 DYGMS 5
      |||||
Db      144 DYGMS 148

RESULT 12
T38737
probable CAAX prenyl proteinase 1 - fission yeast (Schizosaccharomyces pombe)
C;Species: Schizosaccharomyces pombe
C;Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 07-Dec-1999
C;Accession: T38737
R;Gentiles, S.; Churcher, C.M.; Barrell, B.G.; Rajandream, M.A.; Walsh, S.V.
submitted to the EMBL Data Library, December 1995
A;Reference number: Z21808

```

A:Accession: T38737
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-474 <GEN>
A:Cross-references: EMBL:Z68144; PIDN:CAA922258.1; GSPDB:GN00066; SPDB:SPAC3H1.05
A:Experimental source: strain 972h-; cosmid c3H1
C:Genetics:
A:Gene: SPDB:SPAC3H1.05
A:Map position: 1
A:Introns: 35/3

Query Match 100.0%; Score 28; DB 2; Length 474;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 349 DYGMS 353

RESULT 13
F89651
protein T04F8.2 [imported] - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 09-Nov-2001
C:Accession: F89651
R:Anonymous, The C. elegans Sequencing Consortium.
Science 282, 2012-2018, 1998
A:Title: Genome sequence of the nematode C. elegans: a platform for investigating biological processes
A:Reference number: A75000; MUID:99069613; PMID:9851916
A:Note: see websites genome.wustl.edu/gsc/C_elegans/ and www.sanger.ac.uk/projects/C_elegans/
A:Note: published errata appeared in Science 283, 35, 1999; Science 283, 2103, 1999; and Science 283, 2103, 1999; and
A:Accession: F89651
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-478 <STO>
A:Cross-references: GB:chr.X; PIDN:CAA91478.1; PID:G3924836; GSPDB:GN00028; CESP:T04F8.2
A:Note: CDNA EST EMBL:M85094 comes from this gene
C:Genetics:
A:Gene: T04F8.2
A:Map position: X
C:Superfamily: Caenorhabditis elegans hypothetical protein T04F8.2

Query Match 100.0%; Score 28; DB 2; Length 478;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 77 DYGMS 81

RESULT 14
T24471
hypothetical protein T04F8.2 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 04-Mar-2000
C:Accession: T24471
R:Lennard, N.
submitted to the EMBL Data Library, November 1995
A:Reference number: Z19895
A:Accession: T24471
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-502 <WIL>
A:Cross-references: EMBL:Z66565; PIDN:CAA91478.2; GSPDB:GN00028; CESP:T04F8.2
A:Experimental source: clone T04F8
C:Genetics:
A:Gene: CESP:T04F8.2
A:Map position: X
A:Introns: 26/1; 72/1; 103/3; 162/2; 214/3; 249/2; 290/2; 340/3; 382/2; 427/1; 459/2
C:Superfamily: Caenorhabditis elegans hypothetical protein T04F8.2

Query Match 100.0%; Score 28; DB 2; Length 502;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 93 DYGMS 97

RESULT 15
H71285
probable cell division protein (ftsH) - syphilis spirochete
C:Species: Treponema pallidum subsp. pallidum (syphilis spirochete)
C:Date: 24-Jul-1998 #sequence_revision 24-Jul-1998 #text_change 21-Jan-2000
C:Accession: H71285
R:Fraser, C.M.; Norris, S.J.; Weinstock, G.M.; White, O.; Sutton, G.G.; Dodson, R.; Gwin
rson, J.; Khalak, H.; Richardson, D.; Howell, J.K.; Chidambaram, M.; Utterback, T.; McDo
they, L.; Weidman, J.; Smith, H.O.; Venter, J.C.
Science 281, 375-388, 1998
A:Title: Complete genome sequence of Treponema pallidum, the syphilis spirochete.
A:Reference number: A71250; MUID:98332770; PMID:9665876
A:Accession: H71285
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-609 <COL>
A:Cross-references: GB:AE001247; GB:AE000520; NID:G3323059; PIDN:AA065728.1; PID:G332306
A:Experimental source: strain Nichols
C:Genetics:
C:Superfamily: cell division protein ftsH; FtsH/SEC18/CDC48-type ATP-binding domain homo
F:155-363/Domain: FtsH/SEC18/CDC48-type ATP-binding domain homology <VAT>

Query Match 100.0%; Score 28; DB 2; Length 609;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 493 DYGMS 497

Search completed: September 18, 2004, 03:47:32
Job time : 12.1786 secs

FT SITE 57 61 CARBAMOYLPHOSPHATE BINDING (BY
 FT SITE 108 108 CARBAMOYLPHOSPHATE BINDING (BY
 FT SITE 135 135 CARBAMOYLPHOSPHATE BINDING (BY
 FT SITE 148 148 CARBAMOYLPHOSPHATE BINDING (BY
 FT SITE 273 276 IMPORTANT FOR STRUCTURAL INTEGRITY (BY
 FT SITE 334 AA; 37659 MW; BD7A703ABDB63DAE CRC64;
 SQ SEQUENCE 334 AA; 37659 MW; BD7A703ABDB63DAE CRC64;
 Query Match 100.0%; Score 28; DB 1; Length 334;
 Best Local Similarity 100.0%; Pred. No. 34;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QV 1 DYGMS 5
 Db 292 DYGMS 296

RESULT 2

CKR5 MOUSE

ID CKR5 MOUSE STANDARD; PRT; 354 AA.
 AC P51682; O35313; P97308; P97405; Q61867;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE C-C chemokine receptor type 5 (C-C CKR-5) (CCR-5) (MIP-1
 DE alpha receptor).
 GN CKR5 OR CMKERS.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 [1]
 SEQUENCE FROM N.A.
 RC STRAIN=129/SVJ; TISSUE=Spleen;
 RX MEDLINE=96205938; PubMed=8631787;
 RA Boring L., Gosling J., Monteciarlo F.S., Lulis A.J., Tsou C.-L.,
 RA Charo I.F.;
 RT "Molecular cloning and functional expression of murine JE (monocyte
 RT chemoattractant protein 1) and murine macrophage inflammatory protein
 RT 1alpha receptors: evidence for two closely linked C-C chemokine
 RT receptors on chromosome 9.";
 RL J. Biol. Chem. 271:7551-7558(1996).
 [2]
 SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6 X CBA; TISSUE=Thymus;
 RX MEDLINE=96278910; PubMed=8662890;
 RA Meyer A., Coyle A.J., Proudfoot A.E.I., Wells T.N.C., Power C.A.;
 RT "Cloning and characterization of a novel murine macrophage
 RT inflammatory protein-1 alpha receptor.";
 RL J. Biol. Chem. 271:14445-14451(1996).
 [3]
 SEQUENCE FROM N.A.
 RC STRAIN=129/Ola;
 RA Kuziel W.A., Beck M.A., Dawson T.C., Maeda N.;
 RL Submitted (DEC-1996) to the EMBL/GenBank/DBSJ databases.
 [4]
 SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6, and NIH Swiss; TISSUE=Kidney, Liver, and Spleen;
 RX MEDLINE=98001387; PubMed=934322;
 RA Kuhnmann S.E., Platt E.J., Rozak S.L., Kabat D.;
 RT "Polymorphisms in the CKR5 genes of African green monkeys and mice
 RT implicate specific amino acids in infections by simian and human
 RT immunodeficiency viruses.";
 RL J. Virol. 71:8642-8656(1997).
 [5]
 SEQUENCE FROM N.A.
 RC STRAIN=129;
 RX MEDLINE=97404635; PubMed=9261347;
 RA Doranz B.J., Lu Z.H., Rucker J., Zhang T.Y., Sharron M., Cen Y.H.,
 RA Wang Z.X., Guo H.H., Du J.G., Accavitti M.A., Doms R.W., Peiper S.C.;

RT Two distinct CCR5 domains can mediate coreceptor usage by human
 RT immunodeficiency virus type 1.";
 RL J. Virol. 71:6305-6314(1997).
 [6]
 SEQUENCE FROM N.A.
 RA Guo B., Kuno K., Harada A., Matsushima K.;
 RL Submitted (JAN-1997) to the EMBL/GenBank/DBSJ databases.
 CC -1- FUNCTION: Receptor for a C-C type chemokine. Binds to MIP-1-alpha,
 CC MIP-1-beta and RANTES and subsequently transduces a signal by
 CC increasing the intracellular calcium ions level.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- TISSUE SPECIFICITY: Detected in monocyte/macrophage cell lines,
 CC but not in nonhematopoietic cell lines.
 CC -1- SIMILARITY: Belongs to family 1 of G-protein coupled receptors.
 CC
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 CC
 CC EMBL; U47036; AAC52454.1; -
 CC EMBL; X94151; CAA63867.1; -
 CC EMBL; U88585; AAB37273.1; -
 CC EMBL; U83327; AAC53386.1; -
 CC EMBL; AF022990; AAC53389.1; -
 CC EMBL; AF019772; AAB71183.1; -
 CC EMBL; D83648; BAA12024.1; -
 CC MGD; MGI:107182; Ccr5.
 DR GO; GO:0016493; F-C-C chemokine receptor activity; IDA.
 DR GO; GO:0006952; P:defense response; IMP.
 DR InterPro; IPR000276; GPCR_Rhodopsn.
 DR Pfam; PF00001; 7tm_1; 1.
 DR PRINTS; PR00237; GPCRHHODOPSN.
 DR PROSITE; PS00237; G-PROTEIN RECF_F1_1; 1.
 DR PROSITE; PS00362; G-PROTEIN RECF_F1_2; 1.
 KW G-protein coupled receptor; Transmembrane; Glycoprotein; Polymorphism.
 FT DOMAIN 1 32 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 33 60 1 (POTENTIAL).
 FT DOMAIN 61 70 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 71 91 2 (POTENTIAL).
 FT DOMAIN 92 104 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 105 126 3 (POTENTIAL).
 FT DOMAIN 127 143 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 144 168 4 (POTENTIAL).
 FT DOMAIN 169 200 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 201 220 5 (POTENTIAL).
 FT DOMAIN 221 237 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 238 262 6 (POTENTIAL).
 FT DOMAIN 263 279 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 280 303 7 (POTENTIAL).
 FT DOMAIN 304 354 CYTOPLASMIC (POTENTIAL).
 FT DISULFID 103 180 BY SIMILARITY.
 FT CARBOHYD 270 270 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT VARIANT 11 11 I -> S.
 FT VARIANT 62 62 K -> R.
 FT VARIANT 66 66 V -> M.
 FT VARIANT 97 97 I -> V.
 FT VARIANT 109 109 V -> A.
 FT VARIANT 156 156 P -> S.
 FT VARIANT 160 160 F -> L.
 FT VARIANT 185 185 I -> V.
 FT VARIANT 213 213 I -> M.
 FT VARIANT 318 318 V -> A.
 FT VARIANT 337 337 F -> L (IN REF. 2).
 FT CONFLICT 30 30 L -> F (IN REF. 2).
 FT CONFLICT 145 145 N -> I (IN REF. 5).
 FT CONFLICT 190 190 H -> Y (IN REF. 3).
 FT CONFLICT 208 208 P -> S (IN REF. 1).
 FT CONFLICT 354 AA; 40863 MW; B4A6B942E88F9CF0 CRC64;
 SQ SEQUENCE

```
Query Match          100.0%; Score 28; DB 1; Length 354;
Best Local Similarity 100.0%; Pred. No. 36;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Caps 0;

QY 1 DYGMS 5
DB 15 DYGMS 19

RESULT 3
ST24 SCHPO STANDARD; PRT; 474 AA.
AC Q10071;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Probable CAAX prenyl protease 1 (EC 3.4.24.84) (Prenyl protein-
specific endoprotease 1) (PPSEP 1).
GN SPAC3H1.05.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
[1]
RN SEQUENCE FROM N.A.
RC STRAIN=972;
RX MEDLINE=21849401; PubMed=11859360;
RA Wood V., Williams R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgouros J., Peat N., Hayes J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidaigo J., Hodgson G.,
RA Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitsch E.,
RA Rutherford K., Rutter S., Saunders R., Squares S., Stevens K.,
RA Skelton J., Simmonds M., Squires R., Squares S., Sharp S.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volkart G., Aert R., Robben J., Grymonprez B.,
RA Weidjens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritz C., Holzer E., Moestl D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Weller H., Wambutt R., Purnelle B.,
RA Goffeau A., Cadiou E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cerrutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Shpakovski G.V., Ussery D., Barrell B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe."
RL Nature 415:871-880(2002).
CC -!- FUNCTION: Proteolytically removes the C-terminal three residues of
CC -!- farnesylated proteins (By similarity).
CC -!- CATALYTIC ACTIVITY: The peptide bond hydrolyzed can be designated
CC -!- -C[-]RAX in which C is an S-isoprenylated cysteine residue, A is
CC -!- usually aliphatic and X is the C-terminal residue of the substrate
CC -!- protein, and may be any of several amino acids.
CC -!- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Endoplasmic
CC -!- reticulum (By similarity).
CC -!- SIMILARITY: Belongs to peptidase family M48.
CC
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CC EMBL; Z68144; CAA92258.1; -.
DR PIR; T38737; T38737.
DR MEROPS; M48.001; -.
DR GeneDB; Spombe; SPAC3H1.05; -.
DR InterPro; IPR006025; Pept_M_Zn_BS.
DR InterPro; IPR001915; Peptidase_M48.
DR Pfam; PF01435; Peptidase_M48; 1.
DR PROSITE; PS00142; ZINC_PROTEASE; 1.
KW Hypothetical protein; Hydrolase; Metalloprotease; Zinc; Transmembrane;
KW Endoplasmic reticulum.
FT TRANSMEM 103 123 POTENTIAL.
FT TRANSMEM 196 216 POTENTIAL.
FT TRANSMEM 230 250 POTENTIAL.
FT TRANSMEM 344 364 POTENTIAL.
FT TRANSMEM 381 401 POTENTIAL.
FT METAL 332 332 ZINC (CATALYTIC) (BY SIMILARITY).
FT ACT_SITE 333 333 BY SIMILARITY.
FT METAL 336 336 ZINC (CATALYTIC) (BY SIMILARITY).
FT ACT_SITE 411 411 ZINC (CATALYTIC) (BY SIMILARITY).
FT METAL 415 415 PROTON DONOR (BY SIMILARITY).
SQ SEQUENCE 474 AA; 54001 MW; 50AC2F96E25C76C5 CRC64;

Query Match          100.0%; Score 28; DB 1; Length 474;
Best Local Similarity 100.0%; Pred. No. 48;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 349 DYGMS 353

RESULT 4
OCLN_XENLA STANDARD; PRT; 493 AA.
ID OCLN_XENLA STANDARD; PRT; 493 AA.
AC Q9PJN1;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Occludin.
GN OCLN.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
[1]
RN SEQUENCE FROM N.A., AND PHOSPHORYLATION OF THR-375 AND SER-379.
RC TISSUE=Ovary;
RX MEDLINE=99421641; PubMed=10491082;
RA Cordenosi M., Turco F., D'Atri F., Hammar E., Martinucci G.,
RA Meggio F., Citi S.;
RT "Xenopus laevis occludin. Identification of in vitro phosphorylation
sites by protein kinase CK2 and association with cingulin."
RL Eur. J. Biochem. 264:374-384(1999).
RN [2]
RN CHARACTERIZATION.
RX MEDLINE=98034414; PubMed=9365283;
RA Cordenosi M., Mazzone E., De Rigo L., Baraldo S., Meggio F., Citi S.;
RT "Occludin phosphorylation in early development of Xenopus laevis."
RL Cell Sci. 110:3131-3139(1997).
CC -!- FUNCTION: Probably plays a role in the formation and regulation of
CC -!- the tight junction (TJ) paracellular permeability barrier.
CC -!- SUBUNIT: Interacts in vitro with cingulin, possibly directly.
CC -!- Interacts with ZO-1 (By similarity).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
CC -!- TISSUE SPECIFICITY: Localized at tight junctions of both
CC -!- epithelial and endothelial cells.
CC -!- DEVELOPMENTAL STAGE: A maternally synthesized protein. Found in
CC -!- granules in the peripheral cytoplasm in the fertilized egg, it
CC -!- localizes in the peripheral cytoplasm in the basolateral membrane, then to tight
CC -!- junctions after cingulin and ZO-1. Nascent tight junctions are in
CC -!- place by the two-cell stage. The maternal form is more highly
```

phosphorylated than the form detected in later developmental stages.

CC -!- DOMAIN: The C-terminus is cytoplasmic and is important for interaction with ZO-1. Necessary for the tight junction localization. Involved in the regulation of the permeability barrier function of the tight junction (By similarity).

CC -!- PTM: Phosphorylated.

CC -!- SIMILARITY: Belongs to the ELL / occludin family.

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CC -----

DR EMBL; AF170275; AAD53725.1; -.

DR InterPro; IPR008253; MARVEL.

DR InterPro; IPR002958; Occludin.

DR Pfam; PF01284; MARVEL; 1.

DR PRINTS; PR01258; OCCLUDIN.

KW Tight junction; transmembrane; Coiled coil; Phosphorylation.

FT DOMAIN 1 47

FT TRANSMEM 48 70

FT DOMAIN 71 116

FT DOMAIN 117 141

FT DOMAIN 142 151

FT TRANSMEM 152 176

FT DOMAIN 177 224

FT TRANSMEM 225 246

FT DOMAIN 247 493

FT DOMAIN 396 428

FT MOD_RES 375 375

FT MOD_RES 379 379

SQ SEQUENCE 493 AA; 55111 MW; 9694CD302BEBDE CRC64;

Query Match 100.0%; Score 28; DB 1; Length 493;

Best Local Similarity 100.0%; Pred. No. 50;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5

DB 333 DYGMS 337

RESULT 5

FTSH_TREPA STANDARD; PRT; 609 AA.

AC 083746;

DT 15-DEC-1998 (Rel. 37, Created)

DT 15-DEC-1998 (Rel. 37, Last sequence update)

DE 10-OCT-2003 (Rel. 42, Last annotation update)

DE Cell division protein ftsh homolog [EC 3.4.24.-].

GN FTSH OR TPO765.

OS Treponema pallidum.

OC Bacteria; Spirochaetes; Spirochaetales; Spirochaetaceae; Treponema.

OX NCBI_TaxID=160;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=Nichols;

RA Fraser C.M., Norris S.J., Weinstein G.M., White O., Sutton G.G., Dodson R., Gwinn M., Hickey E.K., Clayton R., Ketchum K.A., Sodergren E., Hardham J.M., McLeod M.P., Salzberg S., Peterson J., Khalak H., Richardson D., Howell J.K., Chidambaram M., Utterback T., McDonald L., Artach P., Bowman C., Cotton M.D., Fujii C., Garland S., Hatch B., Horst K., Roberts K., Sandusky M., Weidman J., Smith H.O., Venter J.C.;

RT "Complete genome sequence of Treponema pallidum, the syphilis spirochete."

RL Science 281:375-388 (1998).

CC -!- FUNCTION: Seems to act as an ATP-dependent zinc metallopeptidase

(By similarity).

CC -!- COFACTOR: Binds 1 zinc ion (Potential).

CC -!- SUBCELLULAR LOCATION: Integral membrane protein.

CC -!- SIMILARITY: Belongs to the AAA ATPase family.

CC -!- SIMILARITY: Belongs to peptidase family M41.

CC -----

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CC -----

DR EMBL; AE001247; AAC65728.1; -.

DR TIGR; TP0765; -.

DR InterPro; IPR003593; AAA ATPase.

DR InterPro; IPR003959; AAA_ATPase_cent.

DR InterPro; IPR003960; AAA_sub.

DR InterPro; IPR005936; Peptidase_Ftsh.

DR InterPro; IPR006442; Peptidase_M41.

DR Pfam; PF00004; AAA; 1.

DR Pfam; PF01434; Peptidase_M41; 1.

DR SMART; SM00382; AAA; 1.

DR TIGRFAMs; TIGR01241; Ftsh_fam; 1.

DR PROSITE; PS00674; AAA; 1.

KW Cell division; ATP-binding; Transmembrane; Hydrolase; Metalloprotease; Zinc; Complete proteome.

KW TRANSMEM 89 106

FT NP_BIND 181 188

FT METAL 402 402

FT METAL 403 403

FT ACT_SITE 406 406

FT METAL 406 406

SQ SEQUENCE 609 AA; 67544 MW; 0AE932CA4F6A5D4 CRC64;

Query Match 100.0%; Score 28; DB 1; Length 609;

Best Local Similarity 100.0%; Pred. No. 61;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5

DB 493 DYGMS 497

RESULT 6

PRLR_COLLI STANDARD; PRT; 830 AA.

ID PRLR_COLLI

AC Q90374;

DT 01-NOV-1997 (Rel. 35, Created)

DT 01-NOV-1997 (Rel. 35, Last sequence update)

DT 15-MAR-2004 (Rel. 43, Last annotation update)

DE Prolactin receptor precursor (PRL-R).

GN PRLR.

OS Columba livia (Domestic pigeon).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Archosauria; Aves; Neognathae; Columbiformes; Columba.

OX NCBI_TaxID=8932;

RN [1]

RP SEQUENCE FROM N.A.

RC TISSUE=Cropsac;

RA MEDLINE=94283267; PubMed=7516866;

RT Chen X., Horsman N.D.;

RT "Cloning, expression, and mutational analysis of the pigeon prolactin receptor."

RL Endocrinology 135:269-276 (1994).

CC -!- FUNCTION: This is a receptor for the anterior pituitary hormone prolactin.

CC -!- SUBCELLULAR LOCATION: Type I membrane protein.

CC -!- SIMILARITY: Belongs to the type I cytokine family of receptors. Subfamily 1.

CC -!- SIMILARITY: Contains 4 fibronectin type III domains.

CC -----

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EMBL; U07694; AAA20646.1; --
 PIR; I50455; I50455.
 DR HSSP; P16471; IBP3.
 DR InterPro; IPR002996; CRIA.
 DR InterPro; IPR008957; FN III-like.
 DR InterPro; IPR003961; FN III.
 DR InterPro; IPR003528; Hemtopopn_L_F1.
 DR Pfam; PF00041; fn3; 4.
 DR SMART; SM00060; FN3; 4.
 DR PROSITE; PS01352; HEMATOPO_REC_L_F1; 1.
 KW Receptor; Transmembrane; Glycoprotein; Signal; Repeat.
 FT SIGNAL 1 23
 FT CHAIN 24 830
 FT DOMAIN 24 439
 FT TRANSMEM 440 460
 FT DOMAIN 461 830
 FT DOMAIN 25 122
 FT DOMAIN 123 226
 FT DOMAIN 229 326
 FT DOMAIN 327 429
 FT DISULFID 36 46
 FT DISULFID 75 86
 FT CARBOHYD 59 91
 FT CARBOHYD 91 91
 FT CARBOHYD 100 100
 FT CARBOHYD 112 112
 FT CARBOHYD 132 132
 FT CARBOHYD 263 263
 FT CARBOHYD 304 304
 FT CARBOHYD 316 316
 FT CARBOHYD 336 336
 SQ SEQUENCE 830 AA; 94507 MW; 3B074E83CDF69EFF CRC64;

Query Match 100.0%; Score 28; DB 1; Length 830;
 Best Local Similarity 100.0%; Pred. No. 82;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 77 DYGMS 81

RESULT 7
 ID PLSB_XANAC STANDARD; PRT; 885 AA.
 AC Q8PES0;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Glycerol-3-phosphate acyltransferase (EC 2.3.1.15) (GPAT).
 GN PLSB OR XAC4270.

OS Xanthomonas axonopodis (pv. citri).
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
 OC Xanthomonadaceae; Xanthomonas.
 OX NCBI_TaxID=92829;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=306 / ATCC 13902 / XV 101;
 RX MEDLINE=2022145; PubMed=12024217;
 RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,
 RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
 RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A., L.P.,
 RA Camarotte G., Camnau F., Cardoso J., Chamberg F., Ciapina L.P.,
 RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorry H.,
 RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,

RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
 RA Katsuyana A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
 RA Locali E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
 RA Martins E.C., Medeiros J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
 RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
 RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,
 RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
 RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
 RA Setubal J.C., Kitajima J.P.;
 RT "Comparison of the genomes of two Xanthomonas pathogens with differing
 RT host specificities.";
 RL Nature 417:459-463(2002).
 CC -|- CATALYTIC ACTIVITY: Acyl-CoA + sn-glycerol 3-phosphate = CoA + 1-
 CC acyl-sn-glycerol 3-phosphate.
 CC -|- PATHWAY: De novo phospholipid biosynthesis; first step. May also
 CC function in the regulation of membrane biogenesis.
 CC -|- SUBCELLULAR LOCATION: Membrane-bound (By similarity).
 CC -|- SIMILARITY: Belongs to the GPAT / DAPAT family.

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EMBL; AB012079; AAM39105.1; --
 DR HAMAP; MF 00393; -; 1.
 DR InterPro; IPR002123; Acyltransferase.
 DR Pfam; PF01553; Acyltransferase, 1.
 KW Phospholipid biosynthesis; Transferase; Acyltransferase; Membrane;
 SQ SEQUENCE 885 AA; 98467 MW; FC5942DBAB3B9825 CRC64;

Query Match 100.0%; Score 28; DB 1; Length 885;
 Best Local Similarity 100.0%; Pred. No. 88;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 96 DYGMS 100

RESULT 8
 ID POLN_LORDV STANDARD; PRT; 1699 AA.
 AC P54634;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Non-structural polyprotein [Contains: RNA-directed RNA polymerase
 DE (EC 2.7.7.48); Thiol protease 3C (EC 3.4.22.-); Helicase (2C like
 DE protein)].
 OS Lordsdale virus (Human enteric calicivirus).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Caliciviridae;
 OC Norovirus.
 OX NCBI_TaxID=82658;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=96005060; PubMed=7561776;
 RA Dingle K.E., Lambden P.R., Caul E.O., Clarke I.N.;
 RT "Human enteric Caliciviridae: the complete genome sequence and
 RT expression of virus-like particles from a genetic group II small
 RT round structured virus.";
 RL J. Gen. Virol. 76:2349-2355(1995).
 CC -|- FUNCTION: P2C IS IMPORTANT IN RNA REPLICATION (BY SIMILARITY).
 CC -|- FUNCTION: THE CYSTEINE PROTEASE IS THE PROTEASE RESPONSIBLE FOR
 CC THE POST-TRANSLATIONAL PROCESSING OF THE POLYPEPTIDE (BY
 CC SIMILARITY).
 CC -|- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate +
 CC [RNA](N).
 CC -|- PTM: Specific enzymatic cleavages in vivo yield mature proteins.

CC -!- SIMILARITY: TO PICORNAVIRUS POLYPEPTIDES.
 CC -!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY C24.
 CC
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 CC
 CC EMBL; X86557; CAA60254.1; -
 CC MEROPS; C37.001; -
 CC InterPro; IPR003593; AAA ATPase.
 CC InterPro; IPR004004; Calici pol hel.
 CC InterPro; IPR001665; Peptidase C37.
 CC InterPro; IPR000805; RNA_helicase.
 CC InterPro; IPR007095; RNA_pol_DS_P5.
 CC InterPro; IPR001205; RNA_pol_P3D.
 CC InterPro; IPR007094; RNA_pol_P5vir.
 CC Pfam; PF05416; Peptidase C37; 1.
 CC Pfam; PF06680; RNA_dep_RNA_pol; 1.
 CC Pfam; PF00910; RNA_helicase; 1.
 CC PRINTS; PR00918; CALICIVIRUSNS.
 CC PRINTS; PR00917; SRVCISPTASE.
 CC SMART; SM00382; AAA; 1.
 CC Polyprotein; Transferase; RNA-directed RNA polymerase; Hydrolase;
 KW Thiol protease; Helicase; ATP-binding.
 FT CHAIN ?
 FT CHAIN ?
 FT NP_BIND 495
 FT ACT_SITE 1147
 FT ACT_SITE 1165
 FT ACT_SITE 1165
 SQ SEQUENCE 1699 AA; 189199 MW; FA00B3B67FF3A0B6 CRC64;
 Query Match 100.0%; Score 28; DB 1; Length 1699;
 Best Local Similarity 100.0%; Pred.No. 1.7e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGM5 5
 Db 546 DYGM5 550
 RESULT 9
 ID FAS_FIG STANDARD; PRT; 2133 AA.
 AC P12263; Q95243;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Coagulation factor VIII precursor (Procoagulant component).
 GN F8 OR C8.
 OS Sus scrofa (Pig).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
 OX NCBI_TaxID=9823;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Healey J.F., Lubin I.M., Lollar P.;
 RL Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE OF 705-1573 FROM N.A.
 RX MEDLINE=86287369; PubMed=3016730;
 RA Toole J.J., Pittman D.D., Orr E.C., Murtha P., Wasley L.C.,
 RA Kaufman R.J.;
 RT "A large region (approximately equal to 95 kDa) of human factor VIII
 RT is dispensable for in vitro procoagulant activity."
 RL Proc. Natl. Acad. Sci. U.S.A. 83:5939-5942 (1986).
 RN [3]
 RP SEQUENCE OF 392-759 FROM N.A.
 RX MEDLINE=94179260; PubMed=7510693;
 RA Lubin I.M., Healey J.F., Scandellia D., Runge M.S., Lollar P.;

RT "Elimination of a major inhibitor epitope in factor VIII.";
 J. Biol. Chem. 269:8639-8641 (1994).
 CC -!- FUNCTION: Factor VIII, along with calcium and phospholipid, acts
 CC as a cofactor for factor IXa when it converts factor X to the
 CC activated form, factor Xa.
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- SIMILARITY: Contains 3 F5/8 type A domains.
 CC -!- SIMILARITY: Contains 2 F5/8 type C domains.
 CC -!- SIMILARITY: STRONG, TO COAGULATION FACTOR V.
 CC
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 CC
 CC EMBL; U49517; AAB06705.1; -
 CC PIR; A25945; A25945.
 CC PIR; T42763; T42763.
 CC HSSP; P00451; 1CRG.
 CC InterPro; IPR001117; Cu-oxidase.
 CC InterPro; IPR008972; Cupredoxin.
 CC InterPro; IPR000421; FAS8 C.
 CC InterPro; IPR008979; Gal_Bind_like.
 CC Pfam; PF00394; Cu-oxidase; 3.
 CC Pfam; PF00754; F5_F8 type C; 2.
 CC SMART; SM00231; FAS8C_2.
 CC PROSITE; PS01285; FAS8C_1; 2.
 CC PROSITE; PS01286; FAS8C_2; 2.
 CC PROSITE; PS00022; FAS8C_3; 2.
 CC PROSITE; PS00079; MULTICOPPER_OXIDASE1; 3.
 KW Blood coagulation; Repeat; Plasma; Acute phase; Calcium;
 KW Signal; Glycoprotein; Sulfation.
 FT SIGNAL 1 19 POTENTIAL.
 FT CHAIN 20 2133 COAGULATION FACTOR VIII.
 FT DOMAIN 20 357 F5/8 TYPE A 1.
 FT DOMAIN 20 199 PLASTOCYANIN-LIKE 1.
 FT DOMAIN 207 357 PLASTOCYANIN-LIKE 2.
 FT DOMAIN 399 730 F5/8 TYPE A 2.
 FT DOMAIN 399 573 PLASTOCYANIN-LIKE 3.
 FT DOMAIN 583 730 PLASTOCYANIN-LIKE 4.
 FT DOMAIN 780 1599 B.
 FT DOMAIN 1495 1822 F5/8 TYPE A 3.
 FT DOMAIN 1495 1859 PLASTOCYANIN-LIKE 5.
 FT DOMAIN 1669 1822 PLASTOCYANIN-LIKE 6.
 FT DOMAIN 1822 1970 F5/8 TYPE C 1.
 FT DOMAIN 1975 2127 F5/8 TYPE C 2.
 FT SITE 391 392 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
 FT SITE 759 760 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
 FT SITE 1490 1491 CLEAVAGE (ACTIVATION) (BY SIMILARITY).
 FT SITE 737 737 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
 FT MOD_RES 738 738 SULFATION (BY SIMILARITY).
 FT MOD_RES 742 742 SULFATION (BY SIMILARITY).
 FT DISULFID 173 199 PROBABLE.
 FT DISULFID 547 573 PROBABLE.
 FT DISULFID 1633 1659 PROBABLE.
 FT DISULFID 1822 1970 BY SIMILARITY.
 FT DISULFID 1975 2127 BY SIMILARITY.
 FT CARBOHYD 233 233 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 259 259 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 601 601 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 929 929 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 985 985 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1025 1025 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1111 1111 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1181 1181 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1208 1208 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1245 1245 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1265 1265 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1335 1335 N-LINKED (GLCNAC. . .) (POTENTIAL).

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FT CARBOHYD 1408 1408 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1611 1611 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1919 1919 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 713 713 N -> M (IN REF. 2).
FT CONFLICT 734 734 I -> T (IN REF. 2).
FT CONFLICT 792 792 G -> Q (IN REF. 2).
FT CONFLICT 1133 1133 E -> F (IN REF. 2).
FT CONFLICT 1191 1191 I -> L (IN REF. 2).
FT CONFLICT 1209 1209 R -> F (IN REF. 2).
FT CONFLICT 1437 1437 C -> G (IN REF. 2).
FT CONFLICT 1456 1456 F -> R (IN REF. 2).
FT CONFLICT 1539 1539 F -> R (IN REF. 2).
FT CONFLICT 1546 1546 Q -> N (IN REF. 2).
SQ SEQUENCE 2133 AA; 239304 MW; 152BBA8997F570DA CRC64;

Query Match 100.0%; Score 28; DB 1; Length 2133;
Best Local Similarity 100.0%; Pred. No. 2.1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 1509 DYGMS 1513

RESULT 10
FAS_MOUSE
ID - FAS_MOUSE STANDARD; PRT; 2319 AA.
AC Q06194;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Coagulation factor VIII precursor (Procoagulant component).
GN F8 OR F8C OR F8C.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6 X CBA; TISSUE=Liver;
RA MEDLINE=93300511; PubMed=8314577;
RX Elger B., Laxich D., Gitschier J.;
RT "Sequence of the murine factor VIII cDNA.";
RL Genomics 16:374-379(1993).
CC -!- FUNCTION: Factor VIII, along with calcium and phospholipid, acts
CC as a cofactor for factor IXa when it converts factor X to the
CC activated form, factor Xa.
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- TISSUE SPECIFICITY: Found in most tissues.
CC -!- SIMILARITY: Contains 3 F5/8 type A domains.
CC -!- SIMILARITY: Contains 2 F5/8 type C domains.
CC -!- SIMILARITY: STRONG, TO COAGULATION FACTOR V.
CC
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CC
CC -----
DR EMBL; L05573; AAA37385.1; .
DR F8; A47004; A47004.
DR HSP; P0451; 1CFG.
DR MGD; MGI:88383; F8.
DR InterPro; IPR001117; Cu-oxidase.
DR InterPro; IPR008972; Cupredoxin.
DR InterPro; IPR000421; FAS8_C.
DR Pfam; PF00394; Cu-oxidase; 3.
DR Pfam; PF00754; F5_F8 type_C; 2.
DR SMART; SM00231; FAS8_C2.
DR PROSITE; PS01285; FAS8C_1; 2.
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DR PROSITE; PS01286; FAS8C_2; 2.
DR PROSITE; PS00222; FAS8C_3; 2.
DR PROSITE; PS00079; MULTICOPPER_OXIDASE1; 3.
KW Blood coagulation; Repeat; Plasma; Acute phase; Calcium;
KW Signal; Glycoprotein; Sulfation.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 2319 COAGULATION FACTOR VIII.
FT DOMAIN 20 349 F5/8 TYPE A 1.
FT DOMAIN 20 199 PLASTOCYANIN-LIKE 1.
FT DOMAIN 207 349 PLASTOCYANIN-LIKE 2.
FT DOMAIN 399 730 F5/8 TYPE A 2.
FT DOMAIN 399 573 PLASTOCYANIN-LIKE 3.
FT DOMAIN 583 730 PLASTOCYANIN-LIKE 4.
FT DOMAIN 760 1640 B.
FT DOMAIN 1683 2008 F5/8 TYPE A 3.
FT DOMAIN 1683 1845 PLASTOCYANIN-LIKE 5.
FT DOMAIN 1855 2008 PLASTOCYANIN-LIKE 6.
FT DOMAIN 2008 2156 F5/8 TYPE C 1.
FT DOMAIN 2151 2313 F5/8 TYPE C 2.
FT SITE 391 392 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
FT SITE 759 760 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
FT SITE 1678 1679 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
FT SITE 1324 1325 CLEAVAGE (ACTIVATION) (BY SIMILARITY).
FT SITE 1640 1641 CLEAVAGE (ACTIVATION) (BY SIMILARITY).
FT MOD_RES 367 367 SULFATION (BY SIMILARITY).
FT MOD_RES 737 737 SULFATION (BY SIMILARITY).
FT MOD_RES 738 738 SULFATION (BY SIMILARITY).
FT MOD_RES 742 742 SULFATION (BY SIMILARITY).
FT MOD_RES 1669 1669 SULFATION (REQUIRED FOR VWF BINDING)
(FY SIMILARITY).
FT MOD_RES 1687 1687 SULFATION (BY SIMILARITY).
FT DISULFID 173 199 PROBABLE.
FT DISULFID 547 573 PROBABLE.
FT DISULFID 1839 1845 BY SIMILARITY.
FT DISULFID 2008 2156 BY SIMILARITY.
FT DISULFID 2161 2313 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 61 61 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 233 233 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 259 259 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 423 423 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 601 601 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 880 880 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 958 958 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1015 1015 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1022 1022 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1026 1026 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1044 1044 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1076 1076 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1087 1087 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1136 1136 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1161 1161 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1192 1192 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1255 1255 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1268 1268 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1273 1273 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1274 1274 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1302 1302 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1316 1316 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1340 1340 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1378 1378 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1797 1797 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 2105 2105 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 2319 AA; 266148 MW; FD054DE051DB2A01 CRC64;
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Query Match 100.0%; Score 28; DB 1; Length 2319;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 1697 DYGMS 1701
```

RESULT 11
 FAS_HUMAN STANDARD; PRT; 2351 AA.
 AC P00451;
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Coagulation factor VIII precursor (Procoagulant component)
 DE (Antihemophilic factor) (AHF).
 OS F8 OR F8C.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=86081164; PubMed=3935400;
 RA Truett M.A., Blacher R., Burke R.L., Caput D., Chu C., Dina D.,
 RA Hartog K., Kuo C.H., Masiaz F.R., Merryweather J.P., Najarian R.,
 RA Pacht C., Potter S.J., Puma J., Quiroga M., Rall L.B., Randolph A.,
 RA Urdeda M.S., Valenzuela P., Dahl H.-H.M., Favaloro J., Hansen J.,
 RA Nordfang O., Ezban M.;
 RA "Characterization of the polypeptide composition of human factor
 RT VIII:C and the nucleotide sequence and expression of the human kidney
 RT cDNA.";
 RL DNA 4:333-349(1985).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=85061548; PubMed=6438526;
 RA Wood W.I., Capon D.J., Simonsen C.C., Eaton D.L., Gitschier J.,
 RA Keyt B., Seeburg P.H., Smith D.H., Hollingshead P., Wion K.L.,
 RA Delwart E., Tuddenham E.G.D., Vehar G.A., Lawn R.M.;
 RT "Expression of active human factor VIII from recombinant DNA clones.";
 RL Nature 312:330-337(1984).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=85061550; PubMed=6438528;
 RA Toole J.J., Knopf J.L., Wozney J.M., Sultzman L.A., Buecker J.L.,
 RA Pittman D.D., Kaufman R.J., Brown E., Shoemaker C., Orr E.C.,
 RA Amphlett G.W., Foster W.B., Coe M.L., Knutson G.J., Fass D.N.,
 RA Hewick R.M.;
 RT "Molecular cloning of a cDNA encoding human antihemophilic factor.";
 RL Nature 312:342-347(1984).
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93265012; PubMed=1303178;
 RA Gitschier J., Wood W.I.;
 RT "Sequence of the exon-containing regions of the human factor VIII
 RT gene.";
 RL Hum. Mol. Genet. 1:199-200(1992).
 RN [5]
 RP SEQUENCE OF 2064-2070 FROM N.A.
 RA de Water N.S., Williams R., Browett P.J.;
 RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
 RN [6]
 RP SULFATION OF TYR-1699.
 RX MEDLINE=91093266; PubMed=1898735;
 RA Leyte A., van Schijndel H.B., Niehrs C., Huttner W.B., Verbeet M.P.,
 RA Mertens K., van Mourik J.A.;
 RT "Sulfation of Tyr1680 of human blood coagulation factor VIII is
 RT essential for the interaction of factor VIII with von Willebrand
 RT factor.";
 RL J Biol. Chem. 266:740-746(1991).
 RN [7]
 RP SULFATION.
 RX MEDLINE=92207952; PubMed=1554716;
 RA Pittman D.D., Wang J.H., Kaufman R.J.;
 RT "Identification and functional importance of tyrosine sulfate
 RT residues within recombinant factor VIII.";
 RL Biochemistry 31:3315-3325(1992).
 RN [8]
 RP STRUCTURE BY NMR OF 2322-2343.
 RX MEDLINE=95200924; PubMed=7893714;

RA Gilbert G.E., Baleja J.D.;
 RT "Membrane-binding peptide from the C2 domain of factor VIII forms an
 RT amphipathic structure as determined by NMR spectroscopy.";
 RL Biochemistry 34:3022-3031(1995).
 RN [9]
 RP REVIEW ON MOLECULAR BASIS OF HEMA.
 RX MEDLINE=91221499; PubMed=1902642;
 RA Gitschier J.;
 RT "The molecular basis of hemophilia A.";
 RL Ann. N.Y. Acad. Sci. 614:89-96(1991).
 RN [10]
 RP REVIEW ON MOLECULAR BASIS OF HEMA.
 RX MEDLINE=89088506; PubMed=2491949;
 RA White G.C. II, Shoemaker C.B.;
 RT "Factor VIII gene and hemophilia A.";
 RL Blood 73:1-12(1989).
 RN [11]
 RP REVIEW ON MOLECULAR BASIS OF HEMA.
 RX MEDLINE=95245332; PubMed=7728145;
 RA Antonarakis S.E., Kazazian H.H., Tuddenham E.G.D.;
 RT "Molecular etiology of factor VIII deficiency in hemophilia A.";
 RL Hum. Mutat. 5:1-22(1995).
 RN [12]
 RP VARIANT HEMA GLN-2326.
 RX MEDLINE=86235434; PubMed=3012775;
 RA Gitschier J., Wood W.I., Shuman M.A., Lawn R.M.;
 RT "Identification of a missense mutation in the factor VIII gene of a
 RT mild hemophilic.";
 RL Science 232:1415-1416(1986).
 RN [13]
 RP VARIANT HEMA PRO-2135.
 RX MEDLINE=88096539; PubMed=3122181;
 RA Levinson B., Janco R.L., Phillips J.A. III, Gitschier J.;
 RT "A novel missense mutation in the factor VIII gene identified by
 RT analysis of amplified hemophilia DNA sequences.";
 RL Nucleic Acids Res. 15:9797-9805(1987).
 RN [14]
 RP VARIANT HEMA GLN-2228.
 RX MEDLINE=88191889; PubMed=2833855;
 RA Yousoufian H., Antonarakis S.E., Bell W., Griffin A.M.,
 RA Kazazian H.H.;
 RT "Nonsense and missense mutations in hemophilia A: estimate of the
 RT relative mutation rate at CG dinucleotides.";
 RL Am. J. Hum. Genet. 42:718-725(1988).
 RN [15]
 RP VARIANT HEMA GLY-291.
 RX MEDLINE=88220354; PubMed=2835904;
 RA Yousoufian H., Wong C., Aronis S., Platokoukis H., Kazazian H.H. Jr.,
 RA Antonarakis S.E.;
 RT "Moderately severe hemophilia A resulting from Glu-->Gly substitution
 RT in exon 7 of the factor VIII gene.";
 RL Am. J. Hum. Genet. 42:867-871(1988).
 RN [16]
 RP VARIANT HEMA CYS-1708.
 RX MEDLINE=89274393; PubMed=2499363;
 RA O'Brien D.P., Tuddenham E.G.;
 RT "Purification and characterization of factor VIII 1,689-Cys: a
 RT nonfunctional cofactor occurring in a patient with severe hemophilia
 RT A.";
 RL Blood 73:2117-2122(1989).
 RN [17]
 RP VARIANT HEMA CYS-391.
 RX MEDLINE=90001543; PubMed=2506948;
 RA Shima M., Ware J., Yoshioka A., Fukui H., Fulcher C.A.;
 RT "An arginine to cysteine amino acid substitution at a critical
 RT thrombin cleavage site in a dysfunctional factor VIII molecule.";
 RL Blood 74:1612-1617(1989).
 RN [18]
 RP VARIANT HEMA LEU-189.
 RX MEDLINE=90057680; PubMed=2510835;
 RA Chan V., Chan T.K., Tong T.M., Todd D.;
 RT "A novel missense mutation in exon 4 of the factor VIII:C gene
 RT resulting in moderately severe hemophilia A.";

RL Blood 74:2688-2691(1989).
RN [19]
RP VARIANT HEMA LEU-2326.
RX MEDLINE=89197216; PubMed=2495245;
RA Inaba H., Fujimaki M., Kazazian H.H. Jr., Antonarakis S.E.;
RT "Mild hemophilia A resulting from Arg-to-Leu substitution in exon 26
of the factor VIII gene."
RL Hum. Genet. 81:335-338(1989).
RN [20]
RP VARIANT HEMA HIS-391.
RX MEDLINE=89264502; PubMed=2498882;
RA Arai M., Inaba H., Higuchi M., Antonarakis S.E., Kazazian H.H. Jr.,
RA Fujimaki M., Hoyer L.W.;
RT "Direct characterization of factor VIII in plasma: detection of a
mutation altering a thrombin cleavage site
(arginine-372-->histidine).";
RL Proc. Natl. Acad. Sci. U.S.A. 86:4277-4281(1989).
RN [21]
RP VARIANT HEMA CYS-1708.
RX MEDLINE=90105723; PubMed=2104766;
RA Arai M., Higuchi M., Antonarakis S.E., Kazazian H.H. Jr.,
RA Phillips J.A. III, Janco R.L., Hoyer L.W.;
RT "Characterization of a thrombin cleavage site mutation (Arg 1689 to
Cys) in the factor VIII gene of two unrelated patients with
cross-reacting material-positive hemophilia A."
RL Blood 75:384-389(1990).
RN [22]
RP VARIANTS HEMA GLN-2228 AND LEU-2326.
RX MEDLINE=90123183; PubMed=2105106;
RA Casula L., Murru S., Pecorara M., Ristaldi M.S., Restagno G.,
RA Mancuso G., Morfini M., de Biasi R., Sauto F., Carbonara A.;
RT "Recurrent mutations and three novel rearrangements in the factor
VIII gene of hemophilia A patients of Italian descent."
RL Blood 75:662-670(1990).
RN [23]
RP VARIANT HEMA CYS-391.
RX MEDLINE=90329422; PubMed=1973901;
RA Pattinson J.K., Mcvey J.H., Boon M., Ajani A., Tuddenham E.G.;
RT "CRM+ haemophilia A due to a missense mutation (372-->Cys) at the
internal heavy chain thrombin cleavage site."
RL Br. J. Haematol. 75:73-77(1990).
RN [24]
RP VARIANTS HEMA PHE-1699 AND CYS-1708.
RX MEDLINE=90152591; PubMed=2105306;
RA Higuchi M., Wong C., Kochhan L., Olek K., Aronis S., Kasper C.K.,
RA Kazazian H.H., Antonarakis S.E.;
RT "Characterization of mutations in the factor VIII gene by direct
sequencing of amplified genomic DNA."
RL Genomics 6:65-71(1990).
RN [25]
RP VARIANTS HEMA CYS-1728 AND ASP-1941.
RX MEDLINE=90169898; PubMed=2106480;
RA Traystman M.D., Higuchi M., Kasper C.K., Antonarakis S.E.,
RA Kazazian H.H.;
RT "Use of denaturing gradient gel electrophoresis to detect point
mutations in the factor VIII gene."
Query Match 100.0%; Score 28; DB 1; Length 2351;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 DYGMS 5
DB 1727 DYGMS 1731
RESULT 12
ID RUVX BACSU STANDARD; PRT; 138 AA.
AC O34634.
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Putative Holliday junction resolvase (EC 3.1.-.-).
GN YRRK OR BSU27390.
OC Bacillus subtilis.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
ON NCBI_TaxID=1423;
RX [1]
RP SEQUENCE FROM N.A.
RC STRAIN=168;
RX MEDLINE=98044033; PubMed=9384377;
RA Kunst F., Ogasawara N., Moszer I., Albertini A.M., Alloni G.,
RA Avevedo V., Besterio M.G., Bessieres P., Bolotin A., Borchert S.,
RA Boriss R., Boursier L., Brans A., Braun M., Brignell S.C., Bron S.,
RA Brouillet S., Bruschi C.V., Caldwell B., Capuano V., Carter N.M.,
RA Choi S.K., Codani J.J., Connerthoff A., Cummings N.J., Daniel R.A.,
RA Danizot P., Devine K.M., Dusterhoft A., Ehrlich S.D., Emmerson P.T.,
RA Etian K.D., Errington J., Fabret C., Ferrati E., Foulger D.,
RA Fritz C., Fujita M., Fujita Y., Fuma S., Galizzi A., Galleron N.,
RA Ghim S.Y., Glaser P., Goffeau A., Golligly E.J., Grandi G.,
RA Giuseppe G., Guy B.J., Haga K., Halech J., Harwood C.R., Henaut A.,
RA Hilbert H., Holsappel S., Hosono S., Hullo M.F., Itaya M., Jones L.,
RA Joris B., Karamata D., Kasahara Y., Klaerr-Blanchard M., Klein C.,
RA Kobayashi Y., Koetter P., Koningsstein G., Krogh J., Lazarevic V.,
RA Kuriita K., Lapidus A., Lardinois S., Lauber J., Lazarevic V.,
RA Lee S.M., Levine A., Liu H., Masuda S., Maue C., Medigue C.,
RA Medina N., Mellado R.P., Mizuno M., Moesti D., Nakai S., Noback M.,
RA Noone D., O'Reilly M., Ogawa K., Ogiwara A., Oudega B., Park S.H.,
RA Parro V., Pohl T.M., Portetelle D., Porwollik S., Prescott A.M.,
RA Prescan E., Pujic P., Purnelle B., Rapoport G., Rey M., Reynolds S.,
RA Rieger M., Rivolta C., Rocha E., Roche B., Rose W., Sadaie Y.,
RA Sato T., Scanlan E., Schleich S., Schroeter R., Scoffone F.,
RA Sekiguchi J., Sekowska A., Seror S.J., Serror P., Shin B.S., Soldo B.,
RA Sorokin A., Tacconi E., Takagi T., Takahashi H., Takenaru K.,
RA Takeuchi M., Tamakoshi A., Tanaka T., Terpstra F., Tognoni A.,
RA Tosato V., Uchiyama S., Vandenbol M., Vannier F., Vassarotti A.,
RA Viari A., Wambutt R., Wedler E., Wedler H., Weitzenecker T.,
RA Winters P., Wipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,
RA Yoshida K., Yoshikawa H.F., Zumstein E., Yoshikawa H., Danchin A.;
RT "The complete genome sequence of the Gram-positive bacterium Bacillus
subtilis."
RL Nature 390:249-256(1997).
CC -J- FUNCTION: Could be a nuclease that resolves Holliday junction
intermediates in genetic recombination.
CC -I- SUBCELLULAR LOCATION: Cytoplasmic (Potential).
CC -I- SIMILARITY: Belongs to the YggF HRP family.
CC -----
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CC -----
DR EMBL; Z99117; CAB14681.1; -.
DR PIR; D69979; D69979.
DR Subtilist; BG13792; yrrk.
DR HAMAP; MF_00651; 1.
DR InterPro; IPR005227; Cons_hypoth250.
DR InterPro; IPR006641; YggFc.
DR Pfam; PF03652; UPF0081; 1.
DR SMART; SM00732; YggFc; 1.
DR TIGRFAMs; TIGR00250; TIGR00250; 1.
KW Hydrolase; Nuclease; DNA repair; DNA recombination; Complete proteome.
SQ SEQUENCE 138 AA; 15210 MW; 39F3EA8A63B1C06 CRC64;
Query Match 89.3%; Score 25; DB 1; Length 138;
Best Local Similarity 80.0%; Pred. No. 70;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 DYGMS 5
DB 39 DYGLS 43

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RESULT 13
YMA6_VICFA
ID YMA6_VICFA STANDARD; PRT; 154 AA.
AC Q04655;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical 17.9 kDa protein in AtP6 5' region (ORF154).
OS Vicia faba (Broad bean).
OG Mitochondrion.
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosids I; Fabales; Fabaceae; Papilionoideae; Viciae; Vicia.
OX NCBI_TaxID=3906;
[1]
RN SEQUENCE FROM N.A.
RP MEDLINE=91059426; PubMed=2245478;
RA Macfarlane J.L., Wahlechner J.A., Wolstenholme D.R.;
RT "A broad bean mitochondrial atp6 gene with an unusually simple, non-
RT conserved 5' region";
RL Curr. Genet. 18:87-91(1990).
CC -----
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CC -----
CC EMBL; X54285; CAA38182.1; -.
DR PIR; B61027; B61027.
KW Hypothetical protein; Mitochondrion.
SQ SEQUENCE 154 AA; 17934 MW; 304467D88CC6CE65 CRC64;
Query Match 89.3%; Score 25; DB 1; Length 154;
Best Local Similarity 80.0%; Pred. No. 78;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 DYGMS 5
Db 44 DYGLS 48
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RESULT 14
HASA_SERVA
ID HASA_SERVA STANDARD; PRT; 188 AA.
AC Q54450;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Hemophore HasA (Heme acquisition system protein A).
GN HASA.
OS Serratia marcescens.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Serratia.
OX NCBI_TaxID=615;
[1]
RN SEQUENCE FROM N.A.; SEQUENCE OF 1-11, FUNCTION, SUBCELLULAR LOCATION,
RP AND INDUCTION.
RC STRAIN=SM365;
RX Letoffe S., Ghigo J.M., Wandersman C.;
RT "Iron acquisition from heme and hemoglobin by a Serratia marcescens
RT extracellular protein.";
RL Proc. Natl. Acad. Sci. U.S.A. 91:9876-9880(1994).
[2]
RN SEQUENCE, AND SUBUNIT.
RP MEDLINE=97332469; PubMed=9188703;
RA Izadi N., Henry Y., Haldjian J., Goldberg M.E., Wandersman C.,
RA Delepiere M., Lecroisey A.;
RT "Purification and characterization of an extracellular heme-binding

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RT protein, HasA, involved in heme iron acquisition.";
RL Biochemistry 36:7050-7057(1997).
[3]
RN FUNCTION.
RC STRAIN=SM365;
RX MEDLINE=97315228; PubMed=9171402;
RA Ghigo J.M., Letoffe S., Wandersman C.;
RT "A new type of hemophore-dependent heme acquisition system of Serratia
RT marcescens reconstituted in Escherichia coli.";
RL J. Bacteriol. 179:3572-3579(1997).
[4]
RN X-RAY CRYSTALLOGRAPHY (1.9 ANGSTROMS).
RP MEDLINE=99287096; PubMed=10360351;
RA Arnoux P., Haser R., Izadi N., Lecroisey A., Delepiere M.,
RA Wandersman C., Czjzek M.;
RT "The crystal structure of HasA, a hemophore secreted by Serratia
RT marcescens.";
RL Nat. Struct. Biol. 6:516-520(1999).
[5]
RN X-RAY CRYSTALLOGRAPHY (1.77 ANGSTROMS).
RP MEDLINE=20422928; PubMed=10966573;
RA Arnoux P., Haser R., Izadi-Pruneyre N., Lecroisey A., Czjzek M.;
RT "Functional aspects of the heme bound hemophore HasA by structural
RT analysis of various crystal forms.";
RL Proteins 41:202-210(2000).
CC -!- FUNCTION: Can bind free heme and also acquire it from hemoglobin.
CC Conveys heme from hemoglobin to the HasR receptor which releases
CC it into the bacterium. HasR alone can take up heme but the synergy
CC between HasA and HasR increases heme uptake 100-fold.
CC -!- SUBUNIT: Monomer.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- INDUCTION: By iron depletion.
CC -----
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CC -----
CC EMBL; X81195; CAA57068.1; -.
DR PIR; T48671; T48671.
DR PDB; 1B2V; 24-JUN-99.
DR PDB; 1DKO; 27-DEC-00.
DR PDB; 1DKH; 27-DEC-00.
KW Heme; 3D-structure.
FT METAL 32 32 IRON (HEME AXIAL LIGAND).
FT METAL 75 75 IRON (HEME AXIAL LIGAND).
FT CONFLICT 170 170 A -> Q (IN REF. 2).
SQ SEQUENCE 188 AA; 19283 MW; 8E4ACC8DB254FC6D CRC64;
Query Match 89.3%; Score 25; DB 1; Length 188;
Best Local Similarity 80.0%; Pred. No. 95;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 DYGMS 5
Db 157 DYGLS 161
-----
RESULT 15
EXBB_NEIGO
ID EXBB_NEIGO STANDARD; PRT; 220 AA.
AC Q06433;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Biopolymer transport exbB protein.
GN EXBB.
OS Neisseria gonorrhoeae.
OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
OC Neisseriaceae; Neisseria.

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OK NCBI_TaxID=485;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=FA19;
RX MEDLINE=97285757; PubMed=9140974;
RA Biswas G.D., Anderson J.E., Sparling P.F.;
RT "Cloning and functional characterization of Neisseria gonorrhoeae
RL tonB, exbB and exbD genes.";
CC Mol. Microbiol. 24:169-179(1997).
CC -!- FUNCTION: Involved in the tonB-dependent energy-dependent
CC transport of various receptor-bound substrates. Protects exbD from
CC proteolytic degradation and functionally stabilizes tonB (By
CC similarity).
CC -!- SUBUNIT: The accessory proteins exbB and exbD seem to form a
CC complex with tonB (By similarity).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Inner membrane.
CC -!- SIMILARITY: Belongs to the exbB / tolQ family.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; U79563; AAC45287.1; -.
DR InterPro; IPR002898; Mota_ExbB.
DR Pfam; PF01618; Mota_ExbB; 1.
KW Transport; Protein transport; Transmembrane; Inner membrane.
FT TRANSMEM 13 33 POTENTIAL.
FT TRANSMEM 126 146 POTENTIAL.
FT TRANSMEM 168 188 POTENTIAL.
SQ SEQUENCE 220 AA; 24132 MW; 0247734B36123FBS CRC64;

Query Match 89.3%; Score 25; DB 1; Length 220;
Best Local Similarity 80.0%; Pred. No. 1.1e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
Db 121 DYGMT 125

```

Search completed: September 18, 2004, 03:43:46
Job time : 9.32143 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:24:30 ; Search time 26.0714 Seconds
(without alignments)
60.510 Million cell updates/sec

Title: US-10-029-926b-114
Perfect score: 28
Sequence: 1 DYGMS 5

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SPTREMBL_25.*
1: sp_archaea.*
2: sp_bacteria.*
3: sp_fungi.*
4: sp_human.*
5: sp_invertebrate.*
6: sp_mammal.*
7: sp_mmc.*
8: sp_organelle.*
9: sp_phase.*
10: sp_plant.*
11: sp_rodent.*
12: sp_virus.*
13: sp_vertebrate.*
14: sp_unclassified.*
15: sp_virus.*
16: sp_bacteriap.*
17: sp_archaeap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	70	10 Q84247	Q84247 oryza sativ
2	28	100.0	110	11 Q80Y5	Q80Y5 mus musculu
3	28	100.0	112	4 Q9HCC1	Q9HCC1 homo sapien
4	28	100.0	204	11 Q08826	Q08826 mus musculu
5	28	100.0	204	11 Q08823	Q08823 mus musculu
6	28	100.0	227	16 Q8ZJU0	Q8ZJU0 salmonella
7	28	100.0	227	16 Q8ZUS8	Q8ZUS8 salmonella
8	28	100.0	249	16 Q8E110	Q8E110 streptococc
9	28	100.0	268	10 Q9SKM6	Q9SKM6 arabidopsis
10	28	100.0	268	10 Q81788	Q81788 arabidopsis
11	28	100.0	293	16 Q8PFN0	Q8PFN0 xanthomonas
12	28	100.0	300	10 Q9C840	Q9C840 arabidopsis
13	28	100.0	302	10 Q9MTX8	Q9MTX8 arabidopsis
14	28	100.0	333	11 Q80Y16	Q80Y16 mus musculu
15	28	100.0	340	17 Q9YDP1	Q9YDP1 aeropyrum p
16	28	100.0	362	10 Q9LTC3	Q9LTC3 arabidopsis

17	28	100.0	406	2 Q9EYS9	Q9EYS9 rhizobium s
18	28	100.0	411	16 P72895	P72895 synechocyst
19	28	100.0	439	16 Q87T65	Q87T65 vibrio para
20	28	100.0	448	16 Q880T6	Q880T6 pseudomonas
21	28	100.0	468	16 Q8U8Y0	Q8U8Y0 agrobacteri
22	28	100.0	486	10 Q8L4E4	Q8L4E4 oryza sativ
23	28	100.0	502	5 Q22162	Q22162 caenorhabdi
24	28	100.0	532	10 Q7XR43	Q7XR43 oryza sativ
25	28	100.0	537	12 Q91F86	Q91F86 chilo iride
26	28	100.0	572	3 P87247	P87247 botrytis ci
27	28	100.0	572	16 Q98AA9	Q98AA9 rhizobium l
28	28	100.0	603	16 Q899H3	Q899H3 clostridium
29	28	100.0	650	16 Q8A554	Q8A554 caulobacter
30	28	100.0	728	16 Q9URM7	Q9URM7 rhodospirill
31	28	100.0	756	10 Q9M999	Q9M999 arabidopsis
32	28	100.0	785	4 Q86W01	Q86W01 homo sapien
33	28	100.0	916	10 Q9ZUF5	Q9ZUF5 arabidopsis
34	28	100.0	1008	10 Q9XECO	Q9XECO arabidopsis
35	28	100.0	1076	4 Q8N3A9	Q8N3A9 homo sapien
36	28	100.0	1222	4 Q86T93	Q86T93 homo sapien
37	28	100.0	1236	4 Q96JH7	Q96JH7 homo sapien
38	28	100.0	1250	5 Q23409	Q23409 caenorhabdi
39	28	100.0	1699	12 Q8VOP3	Q8VOP3 human calic
40	28	100.0	1699	12 Q8JX18	Q8JX18 norway-lik
41	28	100.0	1699	12 Q68103	Q68103 hawaii cali
42	28	100.0	1699	12 Q9WI83	Q9WI83 camberwell
43	28	100.0	1699	12 Q8ORD7	Q8ORD7 snow mounta
44	28	100.0	1702	12 Q8CX15	Q8CX15 norway-lik
45	28	100.0	1702	12 Q8CX14	Q8CX14 norway-lik

ALIGNMENTS

RESULT 1

Q84247 ID AC Q84247 PRELIMINARY; PRT; 70 AA.
 DT 01-JUN-2003 (T-EMBLrel. 24, Created)
 DT 01-JUN-2003 (T-EMBLrel. 24, Last sequence update)
 DT 01-JUN-2003 (T-EMBLrel. 24, Last annotation update)
 DE P0458G06.5 protein.
 GN P0458G06.5.
 OS Oryza sativa (japonica cultivar-group).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzoae; Oryza.
 OX NCBI_TaxID=39947;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Nipponbare;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 7, PAC
 clone: P0458G06.5";
 RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AF005051; BAC57333.1; -;
 SQ SEQUENCE 70 AA; 7281 MW; FE752CC92EB2B855 CRC64;

Query Match 100.0%; Score 28; DB 10; Length 70;
 Best Local Similarity 100.0%; Pred. No. 39;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
 |||||
 Db 34 DYGMS 38

RESULT 2

Q80Y5 ID AC Q80Y5 PRELIMINARY; PRT; 110 AA.
 DT 01-JUN-2003 (T-EMBLrel. 24, Created)
 DT 01-JUN-2003 (T-EMBLrel. 24, Last sequence update)

01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 BM289121.4.2 (Novel protein, variant 2) (Fragment).
 BM289121.4.
 Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA North P., Leaves N., Greystrom J., Coppola M., Manjunath S.,
 RA Russell E., Smith M., Strachan G., Tofts C., Boal E., Cobley V.,
 RA Hunter G., Kimberley C., Thomas D., Cave-Berry L., Weston P.,
 RA Botcherby M.R.M.,
 RL Submitted (FEB-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL: BX247953; CAD83028.1; -
 FT NON_TER 1
 SQ SEQUENCE 110 AA; 12302 MW; B5D4F9A90768846F CRC64;
 Query Match 100.0%; Score 28; DB 11; Length 110;
 Best Local Similarity 100.0%; Pred. No. 65;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGMs 5
 DB 49 DYGMs 53

RESULT 3
 Q8HCC1 PRELIMINARY; PRT; 112 AA.
 ID Q8HCC1
 AC Q8HCC1
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Single chain Fv (Fragment).
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Kikuchi M., Takeda C., Tsujimoto Y., Asada S., Nagata K.;
 RA "An antibody fragment2A3 specific for native lysozyme :Isolation from a
 RT human synthetic phage display library and characterization."
 RL Submitted (OCT-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AB049915; BAB16829.1; -
 DR HSSP: P01772; 2FB4.
 DR InterPro: IPR007110; IG-like.
 DR InterPro: IPR003596; IG_v.
 DR Pfam: PF00047; ig; 1.
 DR SMART: SM00406; IGV; 1.
 DR PROSITE: PS50835; IG_LIKE; 1.
 FT NON_TER 1
 FT NON_TER 112
 SQ SEQUENCE 112 AA; 12243 MW; 24F1A45EC3B84788 CRC64;
 Query Match 100.0%; Score 28; DB 4; Length 112;
 Best Local Similarity 100.0%; Pred. No. 67;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGMs 5
 DB 31 DYGMs 35

RESULT 4
 O08826 PRELIMINARY; PRT; 204 AA.
 ID O08826
 AC O08826
 DT 01-JUL-1997 (TrEMBLrel. 04, Created)
 DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE Axonemal dynein heavy chain (Fragment).
 OS Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
 OC Bacterium; Proteobacteria; Gammaproteobacteria; Enterobacteriales;

Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Neesen J., Koehle M., Kirschne R., Steinlein C., Kreutzberger J.,
 RA Engel W., Schmid M.;
 RT "Identification of dynein heavy chain genes expressed in human and
 RT mouse testis: Chromosomal localization of an axonemal dynein gene."
 RL Gene 200:193-202(1997).
 DR EMBL: Z83813; CAB06067.1; -
 FT NON_TER 1
 FT NON_TER 204
 SQ SEQUENCE 204 AA; 22449 MW; 9AFFDA82B4603746 CRC64;
 Query Match 100.0%; Score 28; DB 11; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGMs 5
 DB 199 DYGMs 203

RESULT 5
 O08823 PRELIMINARY; PRT; 204 AA.
 ID O08823
 AC O08823
 DT 01-JUL-1997 (TrEMBLrel. 04, Created)
 DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE Axonemal dynein heavy chain (Fragment).
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Neesen J., Koehle M., Kirschne R., Steinlein C., Kreutzberger J.,
 RA Engel W., Schmid M.;
 RT "Identification of dynein heavy chain genes expressed in human and
 RT mouse testis: Chromosomal localization of an axonemal dynein gene."
 RL Gene 200:193-202(1997).
 DR EMBL: Z83810; CAB06064.1; -
 FT NON_TER 1
 FT NON_TER 204
 SQ SEQUENCE 204 AA; 22198 MW; 72756AFAFB52D0B84 CRC64;
 Query Match 100.0%; Score 28; DB 11; Length 204;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DYGMs 5
 DB 199 DYGMs 203

RESULT 6
 Q8ZJU0 PRELIMINARY; PRT; 227 AA.
 ID Q8ZJU0
 AC Q8ZJU0
 DT 01-MAR-2002 (TrEMBLrel. 20, Created)
 DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Putative fibrial chaparone protein.
 GN STHA OR STW4594.
 OS Salmonella typhimurium.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;

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OC Enterobacteriaceae; Salmonella.
OX NCBI_TaxID=602;
PN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=LT2 / SGSC1412 / ATCC 700720;
RX MEDLINE=21534948; PubMed=11677609;
RA McClelland M., Sanderson K.E., Bentley J., Dougan G., Holden M.T.G., Sebaihia M.,
RA Courtney L., Porwollik S., Ali J., Dante M., Du F., Hou S., Layman D.,
RA Leonard S., Nguyen C., Scott K., Holmes A., Grewal N., Mulvaney E.,
RA Ryan E., Sun H., Flores L., Miller W., Stoneking T., Nhan M.,
RA Waterston R., Wilson R.K.;
RA "Complete genome sequence of Salmonella enterica serovar Typhimurium
RT LT2.";
RL Nature 413:852-856(2001).
RL EMBL; AE008916; AAU23409.1; -.
DR GO; GO:0030288; C:periplasmic space (sensu Gram-negative Bact. . .; IEA.
DR GO; GO:0003754; F:chaperone activity; IEA.
DR GO; GO:0007047; P:cell wall organization and biogenesis; IEA.
DR InterPro; IPR008962; PapD-like.
DR InterPro; IPR001829; Pili_chaperone.
DR Pfam; PF00345; pili_assembly; 1.
DR Pfam; PF02753; pili_assembly; 1.
DR PRINTS; PR00969; CHAPERONPILI.
DR ProDom; PD001447; Pili_chaperone; 1.
DR PROSITE; PS00635; Pili_CHAPERONE; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 227 AA; 24743 MW; EC0D30D9487CD7DF CRC64;

Query Match 100.0%; Score 28; DB 16; Length 227;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 215 DYGMS 219

RESULT 7
ID Q820S8 PRELIMINARY; PRT; 227 AA.
AC Q820S8;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Putative fibrial chaperone protein.
GN STB OR STY493 OR T4633.
OS Salmonella typhi.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Salmonella.
OX NCBI_TaxID=601;
PN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CT18;
RX MEDLINE=21534947; PubMed=11677608;
RA Parkhill J., Dougan G., James K.B., Thomson N.R., Pickard D., Wain J.,
RA Churcher C., Mungall K.L., Bentley S.D., Holden M.T.G., Sebaihia M.,
RA Baker S., Basham D., Brooks K., Chillingworth T., Connor P.,
RA Cronin A., Davis P., Davies R.M., Dowd L., White N., Farrar J.,
RA Feltwell T., Hamlin N., Haque A., Hien T.T., Holroyd S., Jagels K.,
RA Krogh A., Larsen T.S., Leather S., Moule S., O'Garra P., Parry C.,
RA Quail M., Rutherford K., Simmonds M., Skelton J., Stevens K.,
RA Whitehead S., Barrall B.G.;
RA "Complete genome sequence of a multiple drug resistant Salmonella
RT enterica serovar Typhi CT18.";
RL Nature 413:848-852(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=Ty2 / ATCC 700931;
RX MEDLINE=22531367; PubMed=12644504;
RA Deng W., Liou S.-R., Plunkett G. III, Mayhew G.F., Rose D.J.,
RA Burland V., Kodoyanni V., Schwartz D.C., Blattner F.R.;
RT "Comparative genomics of Salmonella enterica serovar Typhi strains Ty2
and CT18.";
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RL J. Bacteriol. 185:2330-2337(2003).
DR EMBL; AL627284; CAD03425.1; -.
DR EMBL; AE016843; AA072062.1; -.
DR GO; GO:0030288; C:periplasmic space (sensu Gram-negative Bact. . .; IEA.
DR GO; GO:0003754; F:chaperone activity; IEA.
DR GO; GO:0007047; P:cell wall organization and biogenesis; IEA.
DR InterPro; IPR008962; PapD-like.
DR InterPro; IPR001829; Pili_chaperone.
DR Pfam; PF00345; pili_assembly; 1.
DR Pfam; PF02753; pili_assembly; 1.
DR PRINTS; PR00969; CHAPERONPILI.
DR ProDom; PD001447; Pili_chaperone; 1.
DR PROSITE; PS00635; Pili_CHAPERONE; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 227 AA; 24829 MW; 6B9F92D7B5421AD1 CRC64;

Query Match 100.0%; Score 28; DB 16; Length 227;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 215 DYGMS 219

RESULT 8
ID Q8E110 PRELIMINARY; PRT; 249 AA.
AC Q8E110;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Prophage LambdaSal, antirepressor, putative.
GN SAG0555.
OS Streptococcus agalactiae (serotype V).
OC Bacteria; Firmicutes; Lactobacillales; Streptococcaceae;
OC Streptococcus.
OX NCBI_TaxID=216466;
PN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=2603 V/R / Serotype V;
RX MEDLINE=2222988; PubMed=12200547;
RA Tetselin H., Masignani V., Cieslewicz M.J., Eisen J.A., Peterson S.,
RA Wessels M.R., Paulsen I.T., Nelson K.E., Margarit I., Read T.D.,
RA Madoff L.C., Wolf A.M., Beanan M.J., Brinkac L.M., Daugherty S.C.,
RA DeBoy R.T., Durkin A.S., Kolonay J.F., Madupu R., Lewis M.R.,
RA Radune D., Fedorova N.B., Scanlan D., Khouri H., Mulligan S.,
RA Carty H.A., Cline R.T., Van Aken S.E., Gill J., Scarselli M., Mora M.,
RA Iacobini E.T., Brettoni C., Galli G., Mariani M., Vegni F., Maione D.,
RA Rinaudo D., Rappuoli R., Telford J.L., Kasper D.L., Grandi G.,
RA Fraser C.M.;
RA "Complete genome sequence and comparative genomic analysis of an
RT emerging human pathogen, serotype V Streptococcus agalactiae.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:12391-12396(2002).
DR EMBL; AE014217; AAM99456.1; -.
DR TIGR; SAG0555; -.
KW Complete proteome.
SQ SEQUENCE 249 AA; 29176 MW; F6537740D2C89258 CRC64;

Query Match 100.0%; Score 28; DB 16; Length 249;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 DYGMS 5
DB 163 DYGMS 167

RESULT 9
ID Q9SKM6 PRELIMINARY; PRT; 268 AA.
AC Q9SKM6;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
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DT 01-MAY-2000 (TReMBLrel. 13, Last sequence update)
 DT 01-OCT-2003 (TReMBLrel. 25, Last annotation update)
 DT En/Spm-like transposon protein.
 GN ATG28440.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 OX NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=20083487; PubMed=10617197;
 RA Lin X., Kaul S., Rounsley S.D., Shea T.P., Benito M.-I., Town C.D.,
 RA Fujii C.Y., Wason T.M., Bowman C.L., Barnstead M.E., Feldblyum T.V.,
 RA Buell C.R., Ketchum K.A., Lee J.J., Ronning C.M., Koo H., Moffat K.S.,
 RA Cronin L.A., Shen M., VanAken S.E., Umayam L., Tallon L.J., Gill J.E.,
 RA Adams M.D., Carrera A.J., Creasy T.H., Goodman H.M., Somerville C.R.,
 RA Salzhager G.P., Preuss D., Nierman W.C., White O., Eisen J.A.,
 RA Salzberg S.L., Fraser C.M., Venter J.C.;
 RT "Sequence and analysis of chromosome 2 of the plant Arabidopsis
 RT thaliana";
 RL Nature 402:761-768(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RA Lin X.;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AC006283; AAD20682.1; -;
 DR PIR: H84684; H84684.
 SQ SEQUENCE 268 AA; 27741 MW; 0584FDE4622FED0E CRC64;

 Query Match 100.0%; Score 28; DB 10; Length 268;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

 QY 1 DYGMS 5
 DB 218 DYGMS 222

 RESULT 10
 CS1798 PRELIMINARY; PRT; 268 AA.
 ID 081788;
 AC 081788;
 DT 01-NOV-1998 (TReMBLrel. 08, Created)
 DT 01-NOV-1998 (TReMBLrel. 08, Last sequence update)
 DT 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
 DE Homeodomain-like protein (AT4G35550/F8D20_60).
 GN F8D20_60 OR AT4G35550.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 OX NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC Koetter P., Hempel S., Entian K.-D., Hoheisel J., Jesse T.,
 RA Heijnen L., Vos P., Mewes H.W., Mayer K.F.X., Schueller C., Bevan M.,
 RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC Rose M., Hempel S., Entian K.-D., Mewes H.W., Lemcke K., Mayer K.F.X.;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC EU Arabidopsis sequencing project;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RC Shinn P., Chen H., Cheuk R., Kim C.J., Koesema E., Meyers M.C.,
 RA Bani J., Bowser L., Carninci P., Dale J.M., Gibson H.A.,
 RA Goldsmith A.D., Hayashizaki Y., Ishida J., Jiang P.X., Jones T.,

RA Kamiya A., Karlin-Neumann G., Kawai J., Lam B., Lee J.M., Lin J.,
 RA Liu S.X., Miranda M., Narusaka M., Nguyen M., Onodera C.S., Palm C.J.,
 RA Pham P.K., Quach H.L., Sakurai T., Satou M., Seki M., Southwick A.,
 RA Tang C.C., Toriumi M., Yamada K., Yu G., Yu S., Shinozaki K.,
 RA Davis R.W., Theologis A., Ecker J.R.;
 RT "Arabidopsis cDNA clones";
 RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
 RN [5]
 RP SEQUENCE FROM N.A.
 RC Cheuk R., Chen H., Kim C.J., Shinn P., Bowser L., Carninci P.,
 RA Chan M.M., Chang C.H., Dale J.M., Deng J.M., Hayashizaki Y.,
 RA Hsuan V.W., Ishida J., Jones T., Kamiya A., Karlin-Neumann G.,
 RA Kawai J., Lam B., Lee J.M., Lin J., Miranda M., Narusaka M.,
 RA Nguyen M., Palm C.J., Quach H.L., Sakurai T., Satou M., Seki M.,
 RA Southwick A., Tang C.C., Toriumi M., Wallender E.K., Wong C., Wu H.C.,
 RA Yamada K., Yu G., Yuan S., Shinozaki K., Davis R.W., Theologis A.,
 Ecker J.R.;
 RT "Arabidopsis ORF clones";
 RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AL031135; CAA20025.1; -;
 DR EMBL: AL161587; CAB80271.1; -;
 DR EMBL: AY048268; AAK82530.1; -;
 DR EMBL: EF000538; AAN18107.1; -;
 DR PIR: T04660; T04660.
 DR GO: GO:0005634; C:nucleus; IEA.
 DR GO: GO:0003700; F:transcription factor activity; IEA.
 DR GO: GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
 DR InterPro: IPR001356; Homeobox.
 DR Pfam: PF00046; homeobox; 1.
 DR Prodom: PD000010; Homeobox; 1.
 DR SMART: SM00389; HOX; 1.
 DR PROSITE: PS0071; HOMEBOX_2; 1.
 KW DNA-binding; Homeobox; Nuclear protein.
 SQ SEQUENCE 268 AA; 29673 MW; D57FCC13AB1A93DA CRC64;

 Query Match 100.0%; Score 28; DB 10; Length 268;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

 QY 1 DYGMS 5
 DB 263 DYGMS 267

 RESULT 11
 CS8PFO PRELIMINARY; PRT; 293 AA.
 ID 08PFO;
 AC 08PFO;
 DT 01-OCT-2002 (TReMBLrel. 22, Created)
 DT 01-OCT-2002 (TReMBLrel. 22, Last sequence update)
 DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
 DE Hypothetical protein XAC3950.
 GN XAC3950.
 OS Xanthomonas axonopodis (pv. citri).
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
 OC Xanthomonadaceae; Xanthomonas.
 OX NCBI_TaxID=92829;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=306 / ATCC 13902 / XV 101;
 RX MEDLINE=2022145; PubMed=12024217;
 RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furian L.R.,
 RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
 RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
 RA Camarotte G., Cannavan F., Cardoso J., Chambergo F., Ciapina L.P.,
 RA Cicarelli R.M.B., Coutinho L.L., Cursino-Santos J.R., El-Dorri H.,
 RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
 RA Formighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
 RA Katsuyama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
 RA Locati E.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
 RA Martins E.C., Meidanis J., Menck C.F.M., Miyaki C.Y., Moon D.H.,
 RA Moreira L.M., Novo M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
 RA Pereira H.A., Rossi A., Sena J.A.D., Silva C., de Souza R.F.,

RA Spinola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
RA Sebail J.C., Kitajima J.P.,
RT "Comparison of the genomes of two Xanthomonas pathogens with differing
RT host specificities",
RL Nature 417:459-463(2002).
DR EMBL; AE012044; AAM38787.1; -
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 293 AA; 33115 MW; 8CAC3C831EA0B33F CRC64;

Query Match 100.0%; Score 28; DB 16; Length 293;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 234 DYGMS 238

RESULT 12

Q9C840 PRELIMINARY; PRT; 300 AA.
AC Q9C840;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN T8524.2.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eucosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RX MEDLINE=21016720; PubMed=11130713;
RA Salanoubat M., Lemcke K., Rieger M., Ansoorge W., Unselid M.,
RA Fartmann B., Valle G., Bloeker H., Perez-Alonso M., Obermaier B.,
RA Delany M., Boutry M., Grivell L.A., Mache R., Puigdomenech P.,
RA De Simone V., Choise N., Artiguenave F., Robert C., Brottier P.,
RA Wincker P., Catolico L., Weissenbach J., Saurin W., Quetier F.,
RA Schaefer M., Mueller-Auer S., Gabel C., Fuchs M., Benes V.,
RA Wurmbach E., Dronek H., Erfle H., Jordan N., Bangert S., Nyakatura G.,
RA Wiedemann R., Kranz H., Voss H., Holland R., Brandt P., Simonati B.,
RA Verzi A., D'Angelo M., Pallavicini A., Toppo S., Simonati B.,
RA Conrad A., Hornischer K., Kauer G., Loehner T.-H., Nordsiek G.,
RA Reichelt J., Scharfe M., Schoen O., Barques M., Terol J., Clement J.,
RA Navarro P., Collado C., Perez-Perez A., Ottenwaelder B., Duchemin D.,
RA Cooke R., Laudie M., Berger-Llauro C., Purnelle B., Masuy D.,
RA de Haan M., Maarse A.C., Alcaraz J.-F., Cottet A., Casacuberta E.,
RA Monfort A., Argirou A., Flores M., Lignori R., Vitale D.,
RA Manhaupt G., Haase D., Schoof H., Rudd S., Zaccaria P., Mewes H.-W.,
RA Mayer K.F.X., Kaul S., Town C.D., Koo H.L., Tallon L.J., Jenkins J.,
RA Rooney T., Rizzo M., Walts A., Utterback T., Fujii C.Y., Shea T.P.,
RA Creasy T.H., Haas B., Maiti R., Wu D., Peterson J., Van Aken S.,
RA Pail G., Mlitscher J., Sellers P., Gill J.E., Feldblyum T.V.,
RA Preuss D., Lin X., Nieman W.C., Salzberg S.L., White O., Venter J.C.,
RA Fraser C.M., Kaneko T., Nakamura Y., Sato S., Kato T., Asamizu E.,
RA Sasamoto S., Kimura T., Idesawa K., Kawashima K., Kishida Y.,
RA Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Muraki A.,
RA Nakayama S., Nakazaki N., Shimpou S., Takeuchi C., Wada T.,
RA Watanabe A., Yamada M., Yasuda M., Tabata S.,
RT "Sequence and analysis of chromosome 3 of the plant Arabidopsis,
RT thaliana.",
RL Nature 408:820-822(2000).
DR EMBL; AC036106; AAG51001.1; -
KW Hypothetical protein.
SQ SEQUENCE 300 AA; 33773 MW; C05A46267AFCE3F4 CRC64;

Query Match 100.0%; Score 28; DB 10; Length 300;
Best Local Similarity 100.0%; Pred. No. 2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 32 DYGMS 36

RESULT 13

Q9M7X8 PRELIMINARY; PRT; 302 AA.
AC Q9M7X8;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE F3E22.15 protein.
GN F3E22.15.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eucosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RA Lin X., Kaul S., Town C.D., Benito M., Creasy T.H., Haas B., Wu D.,
RA Maiti R., Ronning C.M., Koo H., Fujii C.Y., Utterback T.R.,
RA Barnstead M.E., Bowman C.L., White O., Nieman W.C., Fraser C.M.,
RT "Arabidopsis thaliana chromosome III BAC F3E22 genomic sequence.",
RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC023912; AAF63827.1; -
SQ SEQUENCE 302 AA; 34042 MW; C7C97F5F2825AE9 CRC64;

Query Match 100.0%; Score 28; DB 10; Length 302;
Best Local Similarity 100.0%; Pred. No. 2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 32 DYGMS 36

RESULT 14

Q80YV6 PRELIMINARY; PRT; 333 AA.
AC Q80YV6;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE BM289L21.4.1 (Novel protein, variant 1) (Fragment).
GN BM289L21.4.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA North P., Leaves N., Greystrong J., Coppola M., Manjunath S.,
RA Russell E., Smith M., Strachan G., Tofts C., Boal E., Cobley V.,
RA Hunter G., Kimberley C., Thomas D., Cave-Berry L., Weston P.,
RA Botcherby M.R.M.,
RL Submitted (FEB-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BX247953; CAD83027.1; -
FT NON TER 1
SQ SEQUENCE 333 AA; 38205 MW; 42A7BDF256DA90C2 CRC64;

Query Match 100.0%; Score 28; DB 11; Length 333;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DYGMS 5
Db 272 DYGMS 276

```

RESULT 15
QSYDPI
ID QSYDPI PRELIMINARY; PRT; 340 AA.
AC QSYDPI;
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Hypothetical protein APE0874.
GN APE0874.
OS Aeropyrum pernix.
OC Archaea; Crenarchaeota; Thermoprotei; Desulfurococcales;
OC Desulfurococaceae; Aeropyrum.
OX NCBI TaxID=56636;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K1.
RX MEDLINE=9310339; PubMed=10382966;
RA Kwarabayasi Y., Hino Y., Horikawa H., Yamazaki S., Halkawa Y.,
RA Jin-no K., Takahashi M., Sekine M., Baba S.-I., Ankaï A., Kosugi H.,
RA Hosoyama A., Fukui S., Nagai Y., Nishijima K., Nakazawa H.,
RA Takamiya M., Masuda S., Funahashi T., Tanaka T., Kudoh Y.,
RA Yamazaki J., Kishida N., Oguchi A., Aoki K.-I., Kubota K.,
RA Nakamura Y., Nomura N., Sako Y., Kikuchi H.;
RT "Complete genome sequence of an aerobic hyper-thermophilic
RT crenarchaeon, Aeropyrum pernix K1."
RL DNA Res. 6:83-101(1999).
DR EMBL; AP000060; BAA79856.1; -.
DR PIR; H72681; H72681.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 340 AA; 39481 MW; 369F8FA81963C3EF CRC64;

Query Match 100.0%; Score 28; DB 17; Length 340;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 DYGMS 5
Db 82 DYGMS 86

Search completed: September 18, 2004, 03:46:26
Job time : 29.0714 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 02:28:24 ; Search time 120.214 Seconds
(without alignments)
39.956 Million cell updates/sec

Title: US-10-029-926b-115

Perfect score: 96

Sequence: 1 GINWNGSTGYADSVKG 17

Scoring table: BLCSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_29Jan04.*

1: Geneseqp1980s.*

2: Geneseqp1990s.*

3: Geneseqp2000s.*

4: Geneseqp2001s.*

5: Geneseqp2002s.*

6: Geneseqp2003as.*

7: Geneseqp2003bs.*

8: Geneseqp2004s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	DB	ID	Description
1	96	100.0	17	3	AAV95192	AAV95192	Anti-plat
2	96	100.0	17	5	ABG78240	ABG78240	Human Fv
3	96	100.0	17	5	ABG91931	ABG91931	Human ant
4	96	100.0	98	3	AA400073	AA400073	Anti-HiL1
5	96	100.0	98	5	ABG78186	ABG78186	Human Fv
6	96	100.0	98	5	ABG91877	ABG91877	Human ant
7	96	100.0	98	6	ABO27087	ABO27087	Human ger
8	96	100.0	113	3	AAV95177	AAV95177	Anti-plat
9	96	100.0	113	3	AAV95178	AAV95178	Anti-plat
10	96	100.0	115	3	AAV95189	AAV95189	Anti-plat
11	96	100.0	115	3	AAV95190	AAV95190	Anti-plat
12	96	100.0	116	2	AAW19880	AAW19880	CEA-speci
13	96	100.0	117	2	AAW66312	AAW66312	Human imm
14	96	100.0	118	4	AAU02560	AAU02560	Anti-adip
15	96	100.0	122	3	AAV95065	AAV95065	Human ant
16	96	100.0	207	5	AAU98019	AAU98019	Human ace
17	96	100.0	238	3	AAV95198	AAV95198	Anti-plat
18	96	100.0	239	5	ABP46004	ABP46004	Human Bly
19	96	100.0	239	5	ABP46027	ABP46027	Human Bly
20	96	100.0	239	5	ABP44926	ABP44926	Human Bly
21	96	100.0	244	6	AAO31136	AAO31136	Human CM0
22	96	100.0	246	5	ABG78329	ABG78329	Human Fv
23	96	100.0	246	5	ABG93026	ABG93026	Antibody
24	96	100.0	256	5	ABG78334	ABG78334	Human Fv
25	96	100.0	256	5	ABG92025	ABG92025	Antibody

26	96	100.0	266	5	ABG92020	ABG92020	Human ant
27	96	100.0	277	5	ABG78150	ABG78150	Human Fv
28	96	100.0	277	5	ABG78328	ABG78328	Human Fv
29	96	100.0	277	5	ABG92019	ABG92019	Human ant
30	96	100.0	277	5	ABG91841	ABG91841	Human ant
31	96	100.0	309	2	AAW83322	AAW83322	Single ch
32	96	100.0	309	5	ABO9603	ABO9603	Amino aci
33	96	100.0	309	6	ABG74384	ABG74384	Single ch
34	96	100.0	464	5	ABG78151	ABG78151	Human Fv
35	96	100.0	464	5	ABG91842	ABG91842	Human ant
36	93	96.9	126	4	AAU02603	AAU02603	Anti-adip
37	91	94.8	17	3	AA552196	AA552196	Human ant
38	91	94.8	128	3	AA552199	AA552199	Human ant
39	90	93.8	464	5	ABG92021	ABG92021	Antibody
40	79	82.3	98	3	AA340072	AA340072	Anti-HiL1
41	79	82.3	98	5	ABG78185	ABG78185	Human Fv
42	79	82.3	98	5	ABG91876	ABG91876	Human ant
43	79	82.3	99	3	AAV50960	AAV50960	Human Fv
44	79	82.3	99	6	ABO27083	ABO27083	Human ger
45	79	82.3	114	4	AAU02592	AAU02592	Anti-adip

ALIGNMENTS

RESULT 1

AAV95192

ID AAV95192 standard; peptide; 17 AA.

AC AAV95192;

XX

XX 29-AUG-2000 (first entry)

XX

XX Anti-platelet glycoprotein Ib human HIB-1 VH CDR2.

XX

XX Variable heavy chain; single chain antibody; scFv; human; HIB-1;

KW glycoprotein Ib alpha; platelet; aggregation; antiaggregant;

KW antithrombotic; thrombus; therapy; diagnostic; CDR2;

KW complementarity determining region.

XX

XX Homo sapiens.

XX

XX WO200026667-A1.

XX

XX 11-MAY-2000.

XX

XX 29-OCT-1999; 99WO-US025495.

XX

XX 30-OCT-1998; 98US-0106275P.

XX

XX (MILLER) MILLER J L.

XX

XX Miller JL;

XX

XX WPI; 2000-365744/31.

XX

XX Isolated nucleic acid molecule encoding anti-human platelet glycoprotein

PT Ib alpha molecule useful for producing antibodies which inhibit platelet

PT aggregation.

XX

XX Claim 15; Fig 5; 89pp; English.

XX

XX The present sequence is that of complementarity determining region 2

CC (CDR2) of the heavy chain variable region (VH) of human single chain

CC antibody (scFv) HIB-1 (see AAV95198), which is directed against platelet

CC glycoprotein Ib (GPIb). The HIB series of scFv was isolated from a human

CC synthetic VH and VL scFv library on the basis of their binding to

CC platelet GPIb. Whether displayed as surface proteins on a phagemid or

CC secreted as free scFv by Escherichia coli, the HIB scFv clones are

CC capable of inhibiting von Willebrand factor-dependent aggregation of

CC platelets. The scFv are composed of native human protein sequences and

CC are therefore attractive potential reagents for therapeutic purposes.

CC They provide a new class of antithrombotic agents, useful for the

CC prevention of platelet-dependent thrombi in diseased arteries, bypass
 CC grafts, dialysis etc., and can also be used as diagnostic reagents.
 CC Methods of inhibiting aggregation of platelets, of binding human platelet
 CC GPIIb/IIIa and of selecting a VH or VL region of an antibody that
 CC inhibits platelet aggregation are claimed. Fragments of the scFv VH or VL
 CC chain, including CDR fragments, are also claimed

XX Sequence 17 AA;

SQ Query Match 100.0%; Score 96; DB 3; Length 17;
 Best Local Similarity 100.0%; Pred. No. 9.5e-08;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVK 17
 DB 1 GINWNGSGTGADSVK 17

RESULT 2
 ABG78240
 ID ABG78240 standard; peptide; 17 AA.

AC ABG78240;

XX 15-NOV-2002 (first entry)

XX Human Fv molecule hypervariable region related peptide #115.

XX Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
 KW disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
 KW lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.

XX Homo sapiens.

XX WO200259264-A2.

XX 01-AUG-2002.

XX 31-DEC-2001; 2001WO-US049440.

XX 29-DEC-2000; 2000US-00751181.

XX (BIOT-) BIO-TECHNOLOGY GEN CORP.

XX Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
 PI Plaksin D, Peretz T;

XX WPI; 2002-619166/66.

XX Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
 PT or fragment, or construct of fragment with enhanced binding
 PT characteristics so as to selectively bind target cell in favor of other
 PT cells.

PS Claim 20; Page 208-209; 232pp; English.

XX The invention relates to a peptide or polypeptide comprising an Fv
 CC molecule, a construct or fragments or a construct of a fragment with
 CC enhanced binding characteristics which selectively and/or specifically
 CC binds to a target cell in favour of other cells, where binding is
 CC primarily determined by a first hypervariable region and Fv is a single
 CC chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
 CC association with or attached, coupled, combined, linked or fused to a
 CC pharmaceutical agent, is useful in the manufacture of a medicament, where
 CC the medicament has activity against a diseased cell, preferably a cancer
 CC cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
 CC myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
 CC acute myeloid leukaemia cell). The peptide is also useful for preparing a
 CC composition for use in inhibiting the growth of a diseased or cancer
 CC cell. This sequence represents a human Fv molecule hypervariable region
 CC related peptide of the invention

XX Sequence 17 AA;

Query Match 100.0%; Score 96; DB 5; Length 17;
 Best Local Similarity 100.0%; Pred. No. 9.5e-08;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVK 17
 DB 1 GINWNGSGTGADSVK 17

RESULT 3
 ABG91931
 ID ABG91931 standard; peptide; 17 AA.

AC ABG91931;

XX 04-DEC-2002 (first entry)

XX Human antibody fragment #115.

XX Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
 KW metastasis; hypervariable region; autoimmune disease; thrombosis;
 KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
 KW myocardial infarction; retinopathic disease; abnormal platelet function;
 KW sulphated tyrosine-dependent protein-protein interaction.

XX Homo sapiens.

XX WO200253700-A2.

XX 11-JUL-2002.

XX 31-DEC-2001; 2001WO-US049442.

XX 29-DEC-2000; 2000US-00751181.

XX 29-DEC-2000; 2000US-0258948P.

XX (BIOT-) BIO-TECHNOLOGY GEN CORP.

XX Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
 PI Szanton E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;

XX WPI; 2002-674776/72.

XX Novel isolated epitope present on cancer cells and important in
 PT physiological phenomena such as cell rolling, metastasis and
 PT inflammation, for treating autoimmune, inflammatory or cardiovascular
 PT diseases, and cancer.

PS Claim 25; Page 284-285; Opp; English.

XX The invention relates to an isolated epitope present on cancer cells and
 CC important in physiological phenomena such as cell rolling, metastasis and
 CC inflammation, where the epitope is capable of being bound by an antibody,
 CC its antigen-binding fragment or its complex comprising at least one
 CC antibody or its binding fragment having a first hypervariable region. The
 CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
 CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
 CC tumour or leukaemia cells, increase in number of tumour or leukaemia
 CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
 CC platelet and/or cell-platelet adhesion or aggregation, for increasing
 CC mortality of tumour or leukaemia cells, for increasing the susceptibility
 CC of diseased cells to damage by anti-disease, anti-cancer or anti-
 CC leukaemia agents, or for decreasing the number of tumour or leukaemia
 CC cells in a patient, or in the manufacture of a medicament for the above
 CC mentioned purposes. The epitopes are useful for diagnosing and treating
 CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
 CC diseases, cardiovascular diseases such as myocardial infarction,
 CC retinopathic diseases and other diseases mediated by abnormal platelet
 CC function and diseases caused by sulphated tyrosine-dependent protein-
 CC protein interactions. This sequence represents a human antibody fragment
 CC of the invention


```
SQ Sequence 17 AA;
Query Match 100.0%; Score 96; DB 5; Length 17;
Best Local Similarity 100.0%; Pred. No. 9.5e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
DB 1 GINWNGSGTGADSVKG 17

RESULT 4
AAB40073
ID AAB40073 standard; protein; 98 AA.
XX AAB40073;
XX
XX
XX 05-FEB-2001 (first entry)
XX
XX Anti-IL12 antibody H chain V region amino acid sequence SEQ ID 599.
XX Human; neutralising antibody; interleukin-12; IL-12; antiinflammatory;
XX complementarity determining region; CDR; antirheumatic; antiarthritic;
XX antisclerotic; neuroprotective; antipsoaritic; antiasthmatic; cardiant;
XX antiparasitic; antibacterial; immunosuppressive; Crohn's disease;
XX multiple sclerosis; rheumatoid arthritis.
XX
XX Homo sapiens.
XX
XX WO200056772-A1.
XX
XX 28-SEP-2000.
XX
XX 24-MAR-2000; 2000WO-US007946.
XX
XX 25-MAR-1999; 99US-0126603P.
XX
XX (BADI ) BASF AG.
XX (GENY ) GENETICS INST INC.
XX
XX Salfeld JG, Roguska M, Paskind M, Banerjee S, Tracey DE, White M;
XX Kaymakalan Z, Labkovsky B, Sakorafas P, Friedrich S, Myles A;
XX Verdman GM, Venturini A, Warne NW, Widom A, Elvin JG, Duncan AR;
XX Derbyshire EJ, Carmen S, Smith S, Holtet TL, Du Fou SL;
XX WPI; 2000-638250/61.
XX
XX New human antibody specific for human interleukin-12 (IL-12) used to
XX treat disorders characterized by aberrant IL-12 expression e.g. Crohn's
XX disease and multiple sclerosis.
XX
XX Claim 75; Page 121; 377pp; English.
XX
XX This invention relates to a new human antibody specific for human
XX interleukin-12 (IL-12). The invention also includes antigen binding
XX portions that bind to IL-12. Sequences AAB39485-B39516 represent human
XX anti-IL-12 antibody heavy and light chain complementarity determining
XX region (CDR) amino acid sequences, and also includes variable region
XX amino acid sequences. Other variable region amino acid sequences are
XX given in AAB39517-B39560 and AAB40068-B40149. Sequences AAB39561-B39771
XX represent anti-IL-12 CDR3 related amino acid sequences, AAB39772-B40063
XX represent other CDR sequences. Light chain CDR3 consensus sequences are
XX given in AAB40064-B40067. Primers used in the identification and
XX construction of the antibodies of the invention are given in AAC61062-
XX C61071. The antibody of the invention is a neutralising antibody and has
XX antirheumatic; antiarthritic; antisclerotic; antiinflammatory;
XX neuroprotective; antipsoaritic; antiasthmatic; cardiant; antiparasitic;
XX antibacterial and immunosuppressive activity. The antibodies or antigen-
XX binding fragments are useful in the treatment of disorders associated
XX with detrimental release of human IL-12, especially Crohn's disease,
XX multiple sclerosis and rheumatoid arthritis. They can also be used in the
XX manufacture of a pharmaceutical composition to treat human IL-12
XX disorders

XX SQ Sequence 98 AA;
Query Match 100.0%; Score 96; DB 3; Length 98;
Best Local Similarity 100.0%; Pred. No. 6.7e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
DB 50 GINWNGSGTGADSVKG 66

RESULT 5
ABG78186
ID ABG78186 standard; protein; 98 AA.
XX ABG78186;
XX
XX 15-NOV-2002 (first entry)
XX
XX Human Fv molecule hypervariable region related peptide #61.
XX Human; Fv molecule; hypervariable region; single chain Fv; cytostatic;
XX disulfide Fv; dsFv; scFv; cancer; carcinoma; sarcoma; leukaemia; adenoma;
XX lymphoma; myeloma; blastoma; seminoma; melanoma; acute myeloid leukaemia.
XX
XX Homo sapiens.
XX
XX WO200259264-A2.
XX
XX 01-AUG-2002.
XX
XX 31-DEC-2001; 2001WO-US049440.
XX
XX 29-DEC-2000; 2000US-00751181.
XX
XX (BIOT-) BIO-TECHNOLOGY GEN CORP.
XX
XX Hagai Y, Lazarovits J, Guy R, Lipschitz O, Szanton E, Levanon A;
XX Plaksin D, Peretz T;
XX WPI; 2002-619166/66.
XX
XX Novel peptide/polypeptide for cancer therapy has Fv molecule, construct
XX or fragment, or construct of fragment with enhanced binding
XX characteristics so as to selectively bind target cell in favor of other
XX cells.
XX
XX Claim 13; Page 177-178; 232pp; English.
XX
XX The invention relates to a peptide or polypeptide comprising an Fv
XX molecule, a construct or fragments or a construct of a fragment with
XX enhanced binding characteristics which selectively and/or specifically
XX binds to a target cell in favour of other cells, where binding is
XX primarily determined by a first hypervariable region and Fv is a single
XX chain Fv (scFv) or a disulfide Fv (dsFv). The peptide, optionally in
XX association with or attached, coupled, combined, linked or fused to a
XX pharmaceutical agent, is useful in the manufacture of a medicament, where
XX the medicament has activity against a diseased cell, preferably a cancer
XX cell (selected from carcinoma, sarcoma, leukaemia, adenoma, lymphoma,
XX myeloma, blastoma, seminoma, and melanoma, where the leukaemia cell is an
XX acute myeloid leukaemia cell). The peptide is also useful for preparing a
XX composition for use in inhibiting the growth of a diseased or cancer
XX cell. This sequence represents a human Fv molecule hypervariable region
XX related peptide of the invention
XX
XX SQ Sequence 98 AA;
Query Match 100.0%; Score 96; DB 5; Length 98;
Best Local Similarity 100.0%; Pred. No. 6.7e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
```

Db 50 GINWNGSGTGADSVKG 66
|||||
1 GINWNGSGTGADSVKG 17
|||||
50 GINWNGSGTGADSVKG 66

RESULT 6
ABG91877
ID ABG91877 standard; protein; 98 AA.
XX
AC ABG91877;
XX
DT 04-DEC-2002 (first entry)
XX
DE Human antibody fragment #61.
XX
KW Human; antibody; epitope; cancer; tumour; cell rolling; inflammation;
KW metastasis; hypervariable region; autoimmune disease; thrombosis;
KW restenosis; leukaemia; inflammatory disease; cardiovascular disease;
KW myocardial infarction; retinopathic disease; abnormal platelet function;
KW sulphated tyrosine-dependent protein-protein interaction.
XX
OS Homo sapiens.
XX
PN WO200253700-A2.
XX
PD 11-JUL-2002.
XX
PF 31-DEC-2001; 2001WO-US049442.
XX
PR 29-DEC-2000; 2000US-00751181.
PR 29-DEC-2000; 2000US-0258948P.
XX
XX
PA (BIOT-) BIO-TECHNOLOGY GEN CORP.
XX
PI Lazarovits J, Hagai Y, Plaksin D, Vogel T, Nimrod A, Mar-Haim H;
PI Sznathon E, Richter T, Amit B, Kooperman L, Peretz T, Levanon A;
XX
DR WPI; 2002-674776/72.
XX

Novel isolated epitope present on cancer cells and important in
PT physiological phenomena such as cell rolling, metastasis and
PT inflammation, for treating autoimmune, inflammatory or cardiovascular
PT diseases, and cancer.
XX
PS Disclosure; Page 255; Opp; English.
XX

The invention relates to an isolated epitope present on cancer cells and
CC important in physiological phenomena such as cell rolling, metastasis and
CC inflammation, where the epitope is capable of being bound by an antibody,
CC its antigen-binding fragment or its complex comprising at least one
CC antibody or its binding fragment having a first hypervariable region. The
CC epitopes are useful for inhibiting cell rolling, inflammation, autoimmune
CC disease, thrombosis, restenosis, metastasis, growth and/or replication of
CC tumour or leukaemia cells, increase in number of tumour or leukaemia
CC cells in a patient, cell-cell, cell-matrix, platelet-matrix, platelet-
CC platelet and/or cell-platelet adhesion or aggregation, for increasing
CC mortality of tumour or leukaemia cells, for increasing the susceptibility
CC of diseased cells to damage by anti-disease, anti-cancer or anti-
CC leukaemia agents, or for decreasing the number of tumour or leukaemia
CC cells in a patient, or in the manufacture of a medicament for the above
CC mentioned purposes. The epitopes are useful for diagnosing and treating
CC diseases such as cancer, leukaemia, autoimmune diseases, inflammatory
CC diseases, cardiovascular diseases such as myocardial infarction,
CC retinopathic diseases and other diseases mediated by abnormal platelet
CC function and diseases caused by sulphated tyrosine-dependent protein-
CC protein interactions. This sequence represents a human antibody fragment
CC of the invention
XX
SQ Sequence 98 AA;
Query Match 100.0%; Score 96; DB 5; Length 98;
Best Local Similarity 100.0%; Pred. No. 6.7e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 50 GINWNGSGTGADSVKG 66
|||||
1 GINWNGSGTGADSVKG 17
|||||
50 GINWNGSGTGADSVKG 66

RESULT 7
ABO27087
ID ABO27087 standard; protein; 98 AA.
XX
AC ABO27087;
XX
DT 10-SEP-2003 (first entry)
XX
DE Human germline heavy chain variable region gene segment #20.
XX
KW Human; heavy chain variable region; VH; humanised antibody;
KW chimeric antibody; complementarity determining region; CDR;
KW canonical CDR structure type.
XX
OS Homo sapiens.
XX
PN US2003039649-A1.
XX
PD 27-FEB-2003.
XX
PF 12-JUL-2002; 2002US-00194975.
XX
PR 12-JUL-2001; 2001US-0305111P.
XX
PA (FOOT/) FOOTE J.
XX
PI Foote J;
XX
DR WPI; 2003-492151/46.
XX

Making humanized antibody for converting antibody, by making chimeric
PT antibodies containing complementarity determining region from non-human
PT antibody and appropriate framework sequences of human antibodies.
XX
PS Example 1; Fig 1; 31pp; English.
XX

The invention describes a method of making a humanised antibody,
CC comprising making chimeric antibodies containing a complementarity
CC determining region (CDR) from a non-human antibody and appropriate
CC framework sequences (I) of human antibodies. (I) is selected by using
CC canonical CDR structure types of non-human antibody in comparison to
CC germline canonical CDR structure types of human antibodies as the basis
CC for selection, for humanisation. The method is useful for making a
CC humanised antibody or a converted antibody. The method is applicable for
CC converting a subject antibody sequence of any subject species to a less
CC immunogenic form suitable for use in an object species. The method is
CC reliable for identifying suitable human framework sequences to support
CC non-human CDR regions and to provide humanised antibodies that retain
CC high antigen binding with low immunogenicity in humans, without the need
CC for direct comparison of framework sequences, without the need for
CC determining critically important amino acid residues in the framework,
CC and without the need for multiple iteration and construction to obtain
CC humanised antibodies with suitable therapeutic properties. The antibody
CC has high affinity and low immunogenicity without need for comparing
CC framework sequences between non-human and human antibodies. This sequence
CC represents a human heavy chain variable region gene segment used in the
CC creation of humanised antibodies
XX
SQ Sequence 98 AA;
Query Match 100.0%; Score 96; DB 6; Length 98;
Best Local Similarity 100.0%; Pred. No. 6.7e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Query Match 100.0%; Score 96; DB 5; Length 98;
Best Local Similarity 100.0%; Pred. No. 6.7e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

20

CC flooding with murine monoclonal antibody or mimotope peptide (see
 CC AAY95229). Whether displayed as surface proteins on a phagemid or
 CC secreted as free scFv by *Escherichia coli*, the Hib scFv clones are
 CC capable of inhibiting von Willebrand factor-dependent aggregation of
 CC platelets. The scFv are composed of native human protein sequences and
 CC are therefore attractive potential reagents for therapeutic purposes.
 CC They provide a new class of antithrombotic agents, useful for the
 CC prevention of platelet-dependent thrombi in diseased arteries, bypass
 CC grafts, dialysis etc., and can also be used as diagnostic reagents.
 CC Methods of inhibiting aggregation of platelets, of binding human platelet
 CC GPIb alpha and of selecting a VH or VL region of an antibody that
 CC inhibits platelet aggregation are claimed. Note: The present sequence is
 CC not shown in the specification but is derived from the Hib-1 VH sequence
 CC given on page 71 (see AAY95177)
 CC
 XX Sequence 113 AA;
 SQ
 Query Match 100.0%; Score 96; DB 3; Length 113;
 Best Local Similarity 100.0%; Pred. No. 7.8e-07;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GINWNGSGTGADSVKVG 17
 DB 50 GINWNGSGTGADSVKVG 66
 RESULT 10
 AAY95189
 ID AAY95189 standard; protein; 115 AA.
 AC AAY95189;
 XX
 XX 29-AUG-2000 (first entry)
 DT
 DE Anti-platelet glycoprotein Ib human Hib-1 VH.
 XX
 XX Variable heavy chain; single chain antibody; scFv; human; Hib-1;
 KW glycoprotein Ib alpha; platelet; aggregation; antiaggregant;
 KW antithrombotic; thrombus; therapy; diagnostic.
 XX
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT Region 1..30
 FT /note= "framework region 1"
 FT Region 31..34
 FT /note= "complementarity determining region 1"
 FT Region 35..49
 FT /note= "framework region 2"
 FT Region 50..66
 FT /note= "complementarity determining region 2"
 FT Region 67..98
 FT /note= "framework region 3"
 FT Region 99..104
 FT /note= "complementarity determining region 3"
 FT Region 105..115
 FT /note= "framework region 4"
 XX WO200026667-A1.
 FN
 XX 11-MAY-2000.
 PD
 XX 29-OCT-1999; 99WO-US025495.
 XX
 XX 30-OCT-1998; 98US-0106275P.
 PR
 XX (MILLER) MILLER J L.
 PA
 XX Miller JL;
 PI
 XX
 XX WPI; 2000-365744/31.
 DR
 XX Isolated nucleic acid molecule encoding anti-human platelet glycoprotein
 PT

PT Ib alpha molecule useful for producing antibodies which inhibit platelet
 PT aggregation.
 XX
 PS Claim 13; Fig 5; 89pp; English.
 XX
 CC The present sequence is that of the heavy chain variable region (VH) of
 CC human single chain antibody (scFv) Hib-1 (see AAY95198), which is
 CC directed against platelet glycoprotein Ib (GPIb). The Hib series of scFv
 CC was isolated from a human synthetic VH and VL scFv library by 3 rounds of
 CC phagemid selection against transfected CHO cells expressing the GPIb
 CC alpha component of the GPIb/IX/V complex on their surface, followed by a
 CC 4th round of selection against washed human platelets, and 2 final rounds
 CC in which attempts were made to displace scFv from washed platelets by
 CC flooding with murine monoclonal antibody or mimotope peptide (see
 CC AAY95229). Whether displayed as surface proteins on a phagemid or
 CC secreted as free scFv by *Escherichia coli*, the Hib scFv clones are
 CC capable of inhibiting von Willebrand factor-dependent aggregation of
 CC platelets. The scFv are composed of native human protein sequences and
 CC are therefore attractive potential reagents for therapeutic purposes.
 CC They provide a new class of antithrombotic agents, useful for the
 CC prevention of platelet-dependent thrombi in diseased arteries, bypass
 CC grafts, dialysis etc., and can also be used as diagnostic reagents.
 CC Methods of inhibiting aggregation of platelets, of binding human platelet
 CC GPIb alpha and of selecting a VH or VL region of an antibody that
 CC inhibits platelet aggregation are claimed
 CC
 XX Sequence 115 AA;
 SQ

Query Match 100.0%; Score 96; DB 3; Length 115;
 Best Local Similarity 100.0%; Pred. No. 8e-07;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKVG 17
 DB 50 GINWNGSGTGADSVKVG 66

RESULT 11
 AAY95190
 ID AAY95190 standard; protein; 115 AA.
 XX
 XX AAY95190;
 AC
 XX 29-AUG-2000 (first entry)
 DT
 DE Anti-platelet glycoprotein Ib human Hib-1 VH.
 XX
 KW Variable heavy chain; single chain antibody; scFv; human; Hib-1;
 KW glycoprotein Ib alpha; platelet; aggregation; antiaggregant;
 KW antithrombotic; thrombus; therapy; diagnostic.
 XX
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT Region 1..30
 FT /note= "framework region 1"
 FT Region 31..34
 FT /note= "complementarity determining region 1"
 FT Region 35..49
 FT /note= "framework region 2"
 FT Region 50..66
 FT /note= "complementarity determining region 2"
 FT Region 67..98
 FT /note= "framework region 3"
 FT Region 99..104
 FT /note= "complementarity determining region 3"
 FT Region 105..115
 FT /note= "framework region 4"
 XX WO200026667-A1.
 FN
 XX 11-MAY-2000.
 PD
 XX

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PF 29-OCT-1999; 99WO-US025495.
PR 30-OCT-1998; 98US-0106275P.
XX (MILLER) MILLER J L.
XX Miller JL;
XX WPI; 2000-365744/31.
DR WPI; 2000-365744/31.
XX Isolated nucleic acid molecule encoding anti-human platelet glycoprotein
XX Ib alpha molecule useful for producing antibodies which inhibit platelet
XX aggregation.
XX Claim 13; Page; 89pp; English.
XX The present sequence is that of the heavy chain variable region (VH) of
XX human single chain antibody (scFv) H1b-1 (see AA95198), which is
XX directed against platelet glycoprotein Ib (GPIb). The H1b series of scFv
XX was isolated from a human synthetic VH and VL scFv library by 3 rounds of
XX phagemid selection against transfected CHO cells expressing the GPIb
XX alpha component of the GPIb/IX/V complex on their surface, followed by a
XX 4th round of selection against washed human platelets, and 2 final rounds
XX in which attempts were made to displace scFv from washed platelets by
XX flooding with murine monoclonal antibody or mimotope peptide (see
XX AA95529). Whether displayed as surface proteins on a phagemid or
XX secreted as free scFv by Escherichia coli, the H1b scFv clones are
XX capable of inhibiting von Willebrand factor-dependent aggregation of
XX platelets. The scFv are composed of native human protein sequences and
XX are therefore attractive potential reagents for therapeutic purposes.
XX They provide a new class of antithrombotic agents, useful for the
XX prevention of platelet-dependent thrombi in diseased arteries, bypass
XX grafts, dialysis etc., and can also be used as diagnostic reagents.
XX Methods of inhibiting aggregation of platelets, of binding human platelet
XX GPIb alpha and of selecting a VH or VL region of an antibody that
XX inhibits platelet aggregation are claimed. Note: The present sequence is
XX not shown in the specification but is derived from the H1b-1 VH sequence
XX given in Fig 5 (see AA95189)
XX Sequence 115 AA;
SQ
Query Match 100.0%; Score 96; DB 3; Length 115;
Best Local Similarity 100.0%; Pred. No. 8e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db |||||
50 GINWNGSGTGYADSVKG 66

RESULT 12
AAW19880
ID AAW19880 standard; protein; 116 AA.
XX AC AAW19880;
XX 07-DEC-1997 (first entry)
XX CEA-specific antibody CEA5 VH sequence.
XX Carcinoembryonic antigen; CEA; human; antibody; scFv; tumour marker;
XX lung cancer; breast cancer; colon cancer; adenocarcinoma; diagnosis.
XX Homo sapiens.
XX Key Location/Qualifiers
XX Region 31...35
FT /label= CDR1
FT /note= "complementarity determining region 1"
FT Region 50...66
FT /label= CDR2
FT /note= "complementarity determining region 2"
FT Region 99...105

PF 29-OCT-1999; 99WO-US025495.
PR 30-OCT-1998; 98US-0106275P.
XX (MILLER) MILLER J L.
XX Miller JL;
XX WPI; 2000-365744/31.
DR WPI; 2000-365744/31.
XX Isolated nucleic acid molecule encoding anti-human platelet glycoprotein
XX Ib alpha molecule useful for producing antibodies which inhibit platelet
XX aggregation.
XX Claim 13; Page; 89pp; English.
XX The present sequence is that of the heavy chain variable region (VH) of
XX human single chain antibody (scFv) H1b-1 (see AA95198), which is
XX directed against platelet glycoprotein Ib (GPIb). The H1b series of scFv
XX was isolated from a human synthetic VH and VL scFv library by 3 rounds of
XX phagemid selection against transfected CHO cells expressing the GPIb
XX alpha component of the GPIb/IX/V complex on their surface, followed by a
XX 4th round of selection against washed human platelets, and 2 final rounds
XX in which attempts were made to displace scFv from washed platelets by
XX flooding with murine monoclonal antibody or mimotope peptide (see
XX AA95529). Whether displayed as surface proteins on a phagemid or
XX secreted as free scFv by Escherichia coli, the H1b scFv clones are
XX capable of inhibiting von Willebrand factor-dependent aggregation of
XX platelets. The scFv are composed of native human protein sequences and
XX are therefore attractive potential reagents for therapeutic purposes.
XX They provide a new class of antithrombotic agents, useful for the
XX prevention of platelet-dependent thrombi in diseased arteries, bypass
XX grafts, dialysis etc., and can also be used as diagnostic reagents.
XX Methods of inhibiting aggregation of platelets, of binding human platelet
XX GPIb alpha and of selecting a VH or VL region of an antibody that
XX inhibits platelet aggregation are claimed. Note: The present sequence is
XX not shown in the specification but is derived from the H1b-1 VH sequence
XX given in Fig 5 (see AA95189)
XX Sequence 115 AA;
SQ
Query Match 100.0%; Score 96; DB 3; Length 115;
Best Local Similarity 100.0%; Pred. No. 8e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db |||||
50 GINWNGSGTGYADSVKG 66

RESULT 13
AAR66312
ID AAR66312 standard; protein; 117 AA.
XX AC AAR66312;
XX 25-MAR-2003 (revised)
XX 03-AUG-1995 (first entry)
XX Human immunoglobulin variable heavy chain #18.
XX Primer; PCR; amplify; human; immunoglobulin; variable; heavy chain;
XX cosmid; placenta; vector; pJB81; E.coli; mammalian.
XX Homo sapiens.
XX WO9426895-A1.
XX 24-NOV-1994.
XX 10-MAY-1993; 93WO-JP000603.

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FT /label= CDR3
FT /note= "complementarity determining region 3"
XX WO9720932-A1.
XX 12-JUN-1997.
XX 09-DEC-1996; 96WO-GB003043.
XX 07-DEC-1995; 96GB-00025004.
XX 23-MAY-1996; 96GB-00010824.
XX 11-OCT-1996; 96GB-00021295.
XX (CAMP-) CAMBRIDGE ANTIBODY TECHNOLOGY.
XX Osbourn JK, Allen DJ, Mc Cafferty JG;
XX WPI; 1997-319779/29.
XX N-PSDB; AAT72130.
XX Specific binding members for human carcinoembryonic antigen - bind to the
XX A3-B3 extracellular domain of hCEA and are substantially non-cross-
XX reactive with human liver cells; used for diagnosing cancer.
XX Claim 7; Fig 1a; 128pp; English.
XX This polypeptide sequence comprises the heavy chain variable region (VH)
XX of human carcinoembryonic antigen (hCEA)-specific antibody CEA5-VH
XX (AAT72136-32) and VL (AAT72133-35). Gene sequences were obtained for anti-
XX hCEA antibodies CEA1-CEA7 (see AAW19876-85). A claimed specific binding
XX member (A) comprises an hCEA specific antibody antigen binding domain
XX that has a dissociation constant for hCEA of less than 1 x 10-8 M, is
XX non-cross-reactive with human liver cells, and preferentially binds to
XX the A3-B3 extracellular domain of hCEA and/or to cell-associated hCEA
XX over hCEA over soluble hCEA. Preferred (A) include pairings of VH and VL
XX sequences from CEA1-7, or their CDR sequences, as well as CEA5 VH and VL
XX variants. (A) is used to detect cells expressing hCEA, in vivo or in
XX vitro, especially tumour cells for diagnosing cancer, e.g. adenocarcinoma
XX of the colon, lung or breast
XX Sequence 116 AA;
SQ
Query Match 100.0%; Score 96; DB 2; Length 116;
Best Local Similarity 100.0%; Pred. No. 8.1e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db |||||
50 GINWNGSGTGYADSVKG 66

RESULT 13
AAR66312
ID AAR66312 standard; protein; 117 AA.
XX AC AAR66312;
XX 25-MAR-2003 (revised)
XX 03-AUG-1995 (first entry)
XX Human immunoglobulin variable heavy chain #18.
XX Primer; PCR; amplify; human; immunoglobulin; variable; heavy chain;
XX cosmid; placenta; vector; pJB81; E.coli; mammalian.
XX Homo sapiens.
XX WO9426895-A1.
XX 24-NOV-1994.
XX 10-MAY-1993; 93WO-JP000603.

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PR 10-MAY-1993; 93WO-JP000603.
 XX (NIBS) JAPAN TOBACCO INC.
 XX Honjo T, Matsuda F;
 XX WPI; 1995-006791/01.
 DR N-PSDB; AAQ78958.
 XX DNA fragment comprising human immunoglobulin Vh genes - for the
 PT production of human immunoglobulin in mammalian hosts.
 XX
 PS Claim 29; Page 56-57; 130pp; Japanese.
 XX
 CC Protein sequences (AAR66295-51) are novel human immunoglobulin heavy
 CC chain sequences encoded by novel isolated genes. The genes (AAQ78939-
 CC 79002) were isolated and cloned from a series of cosmid constructs: Y202;
 CC Y103; Y21; Y6; Y24; 3-31; M94; M18 and M131, by PCR amplification using
 CC primers AAQ78917-38. The genes are subdivided into 5 families of Vh
 CC genes. The fragments cover a region of 800 kb. The DNA fragments were
 CC isolated from high molecular weight DNA from human placenta. The DNA was
 CC partially digested with TagI restriction enzyme. The fragments were
 CC separated by gel electrophoresis and 35-45 kb fractions were collected.
 CC The fragments were ligated with ClaI-digested cosmid vector pJB81. The
 CC ligation products were in vitro packaged and infected into E.coli 490A. The
 CC fragments were then subcloned by colony hybridisation. The Vh genes and
 CC the DNA fragments encoding them are useful in producing human
 CC immunoglobulin in mammalian hosts. (Updated on 25-MAR-2003 to correct PN
 CC field.)
 XX
 SQ Sequence 117 AA;
 Query Match 100.0%; Score 96; DB 2; Length 117;
 Best Local Similarity 100.0%; Pred. No. 8.2e-07;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GINWNGSGTGADSVK 17
 Db 69 GINWNGSGTGADSVK 85
 RESULT 14
 AAU02560
 ID AAU02560 standard; protein; 118 AA.
 XX
 AC AAU02560;
 XX
 DT 29-AUG-2001 (first entry)
 XX
 DE Anti-adipocyte monoclonal antibody heavy chain, FAT 46.
 XX
 KW Antibody; adipocyte; heavy chain; light chain; obesity; fat;
 XX heart disease; complementarity determining region; CDR.
 XX
 OS Homo sapiens.
 XX
 PN WO200127279-A1.
 XX
 PD 19-APR-2001.
 XX
 PF 11-OCT-2000; 2000WO-GB003900.
 XX
 PR 12-OCT-1999; 99US-0158812P.
 XX
 PA (CAMP-) CAMBRIDGE ANTIBODY TECHNOLOGY.
 XX
 PI Edwards BM, Main SH, Vaughan TJ;
 XX
 DR WPI; 2001-282031/29.
 DR N-PSDB; AAS03460.
 XX
 PT Panel of specific binding members of antibody molecules which bind to
 PT whole adipocytes is used in the treatment of obesity and obesity related

PT diseases.
 XX Claim 1; Page 130; 182pp; English.
 XX
 CC AAU02501-AAU02635, and AAU02641-AAU02748 represent the amino acid
 CC sequences of anti-adipocyte monoclonal antibody heavy chain, light chain,
 CC and heavy chain complementarity determining regions (CDR) of the
 CC invention. The antibodies can be used in the treatment of obesity and
 CC obesity related diseases. The antibodies can be used to deliver drugs or
 CC pro-drugs directly to the fat mass of an obese patient or the antibody
 CC can be used as a therapeutic itself. Antibodies binding specifically to
 CC adipocytes can be used to activate the immune system to destroy the cells
 CC by complement mediated lysis. The antibodies may be labeled with a
 CC detectable label such as radiolabel, fluorescent or chemical group and
 CC used in methods of diagnosis in human subjects e.g. to determine the
 CC presence of adipocyte antigen on the surface of an adipocyte to detect or
 CC determine the presence or level of adipocytes in a cell or tissue sample.
 CC The antibodies can be used as an alternative means of treatment for obese
 CC patients other than undergoing surgery to remove excess fat. Antibodies
 CC for different types of fat deposits can also be produced e.g. intra-
 CC abdominal fat associated with heart disease
 XX
 SQ Sequence 118 AA;
 Query Match 100.0%; Score 96; DB 4; Length 118;
 Best Local Similarity 100.0%; Pred. No. 8.2e-07;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GINWNGSGTGADSVK 17
 Db 50 GINWNGSGTGADSVK 66
 RESULT 15
 AAU02560
 ID AAU02560 standard; protein; 122 AA.
 XX
 AC AAU02560;
 XX
 DT 05-DEC-2000 (first entry)
 XX
 DE Human anti-DAF antibody LU20 heavy chain variable region.
 XX
 KW LU20; human; antibody; VH domain; decay accelerating factor; DAF;
 XX phage display; subtractive panning; lung cancer; lung carcinoma;
 KW lung adenocarcinoma; therapy; diagnosis.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 XX Region 26..32
 FT /note= "hypervariable loop region"
 FT Region 31..35
 FT /label= CDR1
 FT /note= "complementarity determining region I"
 FT Region 50..56
 FT /label= CDRII
 FT /note= "complementarity determining region II"
 FT Region 53..56
 FT /note= "hypervariable loop residues"
 FT Region 99..111
 FT /label= CDRIII
 FT /note= "complementarity determining region III"
 FT Region 100..110
 FT /note= "hypervariable loop residues"
 XX
 PN WO200052054-A2.
 XX
 XX 08-SEP-2000.
 PD
 XX 29-FEB-2000; 2000WO-US0005352.
 PF
 XX 01-MAR-1999; 99US-0122262P.
 PR

XX (GETH) GENENTECH INC.
XX
XX PI Carter PJ, Ridgway JB;
XX
XX DR WPI; 2000-594169/56.
XX
XX PT Making antibodies (e.g. anti-decay accelerating factor antibody) for
XX diagnosing or treating e.g. lung cancer comprises identifying an antigen
XX that is differentially expressed on the surface of two or more distinct
XX cell populations.
XX
XX PS Disclosure; Fig 5B; 52pp; English.
XX
XX CC The present sequence is that of the heavy chain variable region (VH) of
XX the anti-decay accelerating factor (DAF) human antibody LU20. The VL
XX region is given in AAY06062. LU20 was produced using a novel method for
XX making antibodies which can be used for cancer diagnosis or therapy. The
XX method comprises: (a) binding an antibody phase from a naive antibody
XX phage library to a live cancer cell; (b) selecting an antibody phase or
XX antibody which binds selectively to the live cancer cell; and (c)
XX obtaining an antigen to which the antibody phase or antibody binds. To
XX obtain LU20, a human scfv library was used to search for tumour-
XX associated antigens by panning the lung adenocarcinoma cell line 1264,
XX and counter-selecting with a non-tumour bronchial epithelial cell line,
XX BEAS-2B. The invention also describes a method for identifying an antigen
XX which is differentially expressed on the surface of 2 or more distinct
XX cell populations. The anti-DAF human antibody, or a composition
XX comprising the antibody, is useful for in vivo cancer diagnosis or
XX therapy. In particular, the antibody is useful for diagnosing or treating
XX lung cancer, e.g. small-cell lung cancer, non-small cell lung cancer,
XX large cell lung carcinoma, lung adenocarcinoma, or squamous cell lung
XX carcinoma (all claimed)
XX
XX SQ Sequence 122 AA;

Query Match 100.0%; Score 96; DB 3; Length 122;
Best Local Similarity 100.0%; Pred. No. 8.5e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
| | | | | | | | | | | | | | | | | | |
DB 50 GINWNGSGTGYADSVKG 66

Search completed: September 18, 2004, 03:42:54
Job time : 121.214 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:39:30 ; Search time 39.4643 Seconds
(without alignments)
22.239 Million cell updates/sec

Title: US-10-029-926B-115

Perfect score: 96

Sequence: 1 GINWGGSTGYADSVKG 17

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA.*

- 1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
- 2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
- 3: /cgn2_6/ptodata/2/iaa/6A_COMB.pep.*
- 4: /cgn2_6/ptodata/2/iaa/6B_COMB.pep.*
- 5: /cgn2_6/ptodata/2/iaa/PCFUS_COMB.pep.*
- 6: /cgn2_6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	96	100.0	116	2	US-08-652-816A-14
2	96	100.0	117	3	US-08-545-809A-106
3	96	100.0	309	4	US-09-079-029-9
4	79	82.3	118	3	US-08-545-809A-97
5	79	82.3	120	2	US-08-958-201-8
6	79	82.3	120	2	US-08-958-201-10
7	71	74.0	117	3	US-08-545-809A-119
8	71	74.0	118	3	US-08-545-809A-125
9	68	70.8	117	1	US-07-942-245-24
10	63	65.6	126	3	US-08-983-607-26
11	62.5	65.1	117	3	US-09-157-370-1
12	62	64.6	17	1	US-08-264-093-22
13	62	64.6	111	4	US-09-899-896-7
14	62	64.6	118	2	US-08-652-816A-11
15	62	64.6	120	1	US-08-264-093-14
16	62	64.6	483	3	US-09-049-672A-5
17	61.5	64.1	95	3	US-09-043-514-2
18	61.5	64.1	116	3	US-08-974-899-6
19	61	63.5	113	3	US-08-545-809A-135
20	61	63.5	116	3	US-09-184-658-48
21	61	63.5	116	3	US-08-983-607-36
22	61	63.5	116	4	US-09-504-262D-48
23	61	63.5	117	3	US-08-545-809A-103
24	61	63.5	117	3	US-08-545-809A-109
25	61	63.5	117	3	US-08-983-607-46
26	61	63.5	118	2	US-08-652-816A-12
27	61	63.5	119	4	US-09-648-067A-15

28	61	63.5	120	2	US-08-428-197-20	Sequence 20, Appl
29	61	63.5	120	2	US-08-428-197-22	Sequence 22, Appl
30	61	63.5	120	2	US-08-428-197-24	Sequence 24, Appl
31	61	63.5	120	2	US-08-428-197-26	Sequence 26, Appl
32	61	63.5	120	2	US-08-428-197-28	Sequence 28, Appl
33	61	63.5	120	2	US-08-428-197-30	Sequence 30, Appl
34	61	63.5	120	2	US-08-428-197-32	Sequence 32, Appl
35	61	63.5	120	2	US-08-428-197-34	Sequence 34, Appl
36	61	63.5	120	2	US-08-428-197-40	Sequence 40, Appl
37	61	63.5	120	4	US-09-025-769B-38	Sequence 38, Appl
38	61	63.5	120	4	US-09-025-769B-63	Sequence 63, Appl
39	61	63.5	120	5	PCT-US93-07832-4	Sequence 4, Appl
40	61	63.5	120	5	PCT-US93-10555-20	Sequence 20, Appl
41	61	63.5	120	5	PCT-US93-10555-22	Sequence 22, Appl
42	61	63.5	120	5	PCT-US93-10555-24	Sequence 24, Appl
43	61	63.5	120	5	PCT-US93-10555-26	Sequence 26, Appl
44	61	63.5	120	5	PCT-US93-10555-28	Sequence 28, Appl
45	61	63.5	120	5	PCT-US93-10555-30	Sequence 30, Appl

ALIGNMENTS

RESULT 1
US-08-652-816A-14
; Sequence 14, Application US/08652816A
; Patent No. 5872215
; GENERAL INFORMATION:
; APPLICANT: Osbourn, JK
; APPLICANT: Allen, DJ
; APPLICANT: McCafferty, JG
; TITLE OF INVENTION: Specific binding members, materials and
; TITLE OF INVENTION: methods.
; NUMBER OF SEQUENCES: 53
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; COMPUTER READABLE FORM: disk
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (SPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,816A
; FILING DATE: 23-MAY-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9125579.4
; FILING DATE: 02-DEC-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9125579.8
; FILING DATE: 02-DEC-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9206318.9
; FILING DATE: 24-MAR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9206372.6
; FILING DATE: 23-SEP-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9525004.9
; FILING DATE: 07-DEC-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9610824.6
; FILING DATE: 23-MAY-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB92/02240
; FILING DATE: 02-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/244,597
; FILING DATE: 01-JUN-1994
; ATTORNEY/AGENT INFORMATION:

NAME: David W. Clough
 REGISTRATION NUMBER: 36,107
 REFERENCE/DOCKET NUMBER: 28111/33308
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 312-474-6300
 INFORMATION FOR SEQ ID NO: 14:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 116 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 US-08-652-816A-14

Query Match 100.0%; Score 96; DB 2; Length 116;
 Best Local Similarity 100.0%; Pred. No. 1.7e-06;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
 DB 50 GINWNGSGTGADSVKG 66

RESULT 2

US-08-545-809A-106
 Sequence 106, Application US/08545809A
 Patent No. 6096878

GENERAL INFORMATION:
 APPLICANT: Honjo, Tasuku
 APPLICANT: Matsuda, Fumihiko
 TITLE OF INVENTION: HUMAN IMMUNOGLOBULIN VH GENE
 NUMBER OF SEQUENCES: 145
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Fish & Richardson, P.C.
 STREET: 225 Franklin Street
 CITY: Boston
 STATE: MA
 COUNTRY: US
 ZIP: 02110-2804

COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: Windows95
 SOFTWARE: FastSeq for Windows Version 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/545.809A
 FILING DATE: 27-MAR-1996

PRIOR APPLICATION DATA:
 APPLICATION NUMBER: PCT/JP93/00603
 FILING DATE: 10-MAY-1993
 ATTORNEY/AGENT INFORMATION:
 NAME: Freeman, John W.
 REGISTRATION NUMBER: 29,066
 REFERENCE/DOCKET NUMBER: 08501/004001
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 617-542-5070
 TELEFAX: 617-542-8906
 TELEX: 200154

INFORMATION FOR SEQ ID NO: 106:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 117 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-545-809A-106

Query Match 100.0%; Score 96; DB 3; Length 117;
 Best Local Similarity 100.0%; Pred. No. 1.7e-06;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
 DB 69 GINWNGSGTGADSVKG 85

RESULT 3

US-09-079-029-9
 Sequence 9, Application US/09079029
 Patent No. 6342369

GENERAL INFORMATION:
 APPLICANT: Adams, Camilia W.
 APPLICANT: Ashkenazi, Avi J.
 APPLICANT: Chuntharapai, Anan
 APPLICANT: Kim, Kyung J.
 TITLE OF INVENTION: Apo-2 Receptor
 NUMBER OF SEQUENCES: 14
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Genentech, Inc.
 STREET: 1 DNA Way
 CITY: South San Francisco
 STATE: California
 COUNTRY: USA
 ZIP: 94080

COMPUTER READABLE FORM:
 MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: WinPatIn (Genentech)
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/079,029

FILING DATE:

CLASSIFICATION:
 ATTORNEY/AGENT INFORMATION:
 NAME: Marschang, Diane L.
 REGISTRATION NUMBER: 35,600
 REFERENCE/DOCKET NUMBER: P1101R2
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 650/225-5416
 TELEFAX: 650/952-9881
 INFORMATION FOR SEQ ID NO: 9:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 309 amino acids
 TYPE: Amino Acid
 TOPOLOGY: Linear
 US-09-079-029-9

Query Match 100.0%; Score 96; DB 4; Length 309;
 Best Local Similarity 100.0%; Pred. No. 4.7e-06;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
 DB 89 GINWNGSGTGADSVKG 105

RESULT 4

US-08-545-809A-97
 Sequence 97, Application US/08545809A
 Patent No. 6096878

GENERAL INFORMATION:
 APPLICANT: Honjo, Tasuku
 APPLICANT: Matsuda, Fumihiko
 TITLE OF INVENTION: HUMAN IMMUNOGLOBULIN VH GENE
 NUMBER OF SEQUENCES: 145
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Fish & Richardson, P.C.
 STREET: 225 Franklin Street
 CITY: Boston
 STATE: MA
 COUNTRY: US
 ZIP: 02110-2804

COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: Windows95
 SOFTWARE: FastSeq for Windows Version 2.0

SEQUENTS AND DNA FRAGMENTS CONTAINING THE SAME

;;
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/545.809A
;; FILING DATE: 27-MAR-1996
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: PCT/JP93/00603
;; FILING DATE: 10-MAY-1993
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Freeman, John W.
;; REGISTRATION NUMBER: 29,066
;; REFERENCE/DOCKET NUMBER: 06501/004001
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 617-542-5070
;; TELEFAX: 617-542-8906
;; TELEX: 200154
;; INFORMATION FOR SEQ ID NO: 97:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 118 amino acids
;; TYPE: amino acid
;; TOPOLOGY: linear
;; MOLECULE TYPE: protein
US-08-545-809A-97

Query Match 82.3%; Score 79; DB 3; Length 118;
Best Local Similarity 82.4%; Pred. No. 0.00043;
Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db 69 GISWNSGSGYADSVKG 85
||:|||||
||:|||||

RESULT 5
US-08-958-201-8
; Sequence 8, Application US/08958201
; Patent No. 5977319
; GENERAL INFORMATION:
; APPLICANT: Pope, Anthony R.
; APPLICANT: Pritchard, Kevin J
; APPLICANT: Williams, Andrew J
; APPLICANT: Johnson, Kevin S
; TITLE OF INVENTION: Specific binding members for estradiol;
; TITLE OF INVENTION: materials and methods
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall O'Toole Gerstein Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606-6402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.25 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/958,201
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/028,897
; FILING DATE: 21-OCT-1996
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 120 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 2DB
US-08-958-201-10

Query Match 82.3%; Score 79; DB 2; Length 120;
Best Local Similarity 82.4%; Pred. No. 0.00044;
Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db 50 GISWNSGSGYADSVKG 66
||:|||||
||:|||||

RESULT 7
US-08-545-809A-119
; Sequence 119, Application US/08545809A
; Patent No. 6096878
; GENERAL INFORMATION:
; APPLICANT: Honjo, Tasuku
; APPLICANT: Matsuda, Fumihiko
; TITLE OF INVENTION: HUMAN IMMUNOGLOBULIN VH GENE
; TITLE OF INVENTION: SEGMENTS AND DNA FRAGMENTS CONTAINING THE SAME
; NUMBER OF SEQUENCES: 145
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson, P.C.
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: MA
; COUNTRY: US
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette

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; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows95
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/545,809A
; FILING DATE: 27-MAR-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/JP93/00603
; FILING DATE: 10-MAY-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Freeman, John W.
; REGISTRATION NUMBER: 29,066
; REFERENCE/DOCKET NUMBER: 06501/004001
; TELEPHONE: 617-542-5070
; TELEFAX: 617-542-8906
; TELEX: 200154
; INFORMATION FOR SEQ ID NO: 119:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-545-809A-119

Query Match      74.0%; Score 71; DB 3; Length 117;
Best Local Similarity 70.6%; Pred. No. 0.0056;
Matches 12; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY      1 GINWNGSGTGYADSVKG 17
      |:|||||:|||||
Db      59 GVSNGSRTHYADSVKG 85

RESULT 8
US-08-545-809A-125
; Sequence 125, Application US/08545809A
; Patent No. 6096878
; GENERAL INFORMATION:
; APPLICANT: Matsuda, Fumihiko
; APPLICANT: Honjo, Tasuku
; TITLE OF INVENTION: HUMAN IMMUNOGLOBULIN VH GENE
; TITLE OF INVENTION: SEGMENTS AND DNA FRAGMENTS CONTAINING THE SAME
; NUMBER OF SEQUENCES: 145
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson, P.C.
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: MA
; COUNTRY: US
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; OPERATING SYSTEM: Windows95
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/545,809A
; FILING DATE: 27-MAR-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/JP93/00603
; FILING DATE: 10-MAY-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Freeman, John W.
; REGISTRATION NUMBER: 29,066
; REFERENCE/DOCKET NUMBER: 06501/004001
; TELEPHONE: 617-542-5070
; TELEFAX: 617-542-8906
; TELEX: 200154
; INFORMATION FOR SEQ ID NO: 125:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 118 amino acids
```

```
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-545-809A-125

Query Match      74.0%; Score 71; DB 3; Length 118;
Best Local Similarity 81.2%; Pred. No. 0.0057;
Matches 13; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY      2 INWNGSGTGYADSVKG 17
      |:|||||:|||||
Db      70 ISWDGSGTYADSVKG 85

RESULT 9
US-07-942-245-24
; Sequence 24, Application US/079422245
; Patent No. 5639641
; GENERAL INFORMATION:
; APPLICANT: PEDERSEN, Jan T.
; APPLICANT: SEARLE, Stephen M.J.
; APPLICANT: REES, Anthony R.
; APPLICANT: ROGUSKA, Michael A.
; APPLICANT: GUILD, Braydon C.
; TITLE OF INVENTION: SURFACE RESIDUE VENEERING OF RODENT
; TITLE OF INVENTION: ANTIBODIES
; NUMBER OF SEQUENCES: 522
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sughrue, Mion, Zinn, Macpeak & Seas
; STREET: 2100 Pennsylvania Avenue, N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: United States
; ZIP: 20037-3202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: HP 9000/700 Workstation
; OPERATING SYSTEM: UNIX
; SOFTWARE: in house
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/942,245
; FILING DATE: 09-SEP-1992
; CLASSIFICATION: 530
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 293-7060
; TELEFAX: (202) 293-7860
; TELEX: 6491103
; INFORMATION FOR SEQ ID NO: 24:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-07-942-245-24

Query Match      70.8%; Score 68; DB 1; Length 117;
Best Local Similarity 70.6%; Pred. No. 0.015;
Matches 12; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY      1 GINWNGSGTGYADSVKG 17
      |:|||||:|||||
Db      50 GISWDSSIGYADSVKG 66

RESULT 10
US-08-983-607-26
; Sequence 26, Application US/08983607
; Patent No. 6140470
; GENERAL INFORMATION:
; APPLICANT: Alan Garen
; APPLICANT: Xiaohong Cai
; TITLE OF INVENTION: Human Anti-Tumor Monoclonal Anti-
; TITLE OF INVENTION: bodies
```

NUMBER OF SEQUENCES: 51
CORRESPONDENCE ADDRESS:
ADDRESSEE: Department of Molecular Biophysics
ADDRESSEE: and Biochemistry, Yale University
STREET: 266 Whitney Avenue
CITY: New Haven
STATE: Connecticut
COUNTRY: United States of America
ZIP: 06520-8114
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" 1.44 Mb diskette
COMPUTER: IBM PC
OPERATING SYSTEM: MS DOS
SOFTWARE: Word Processing
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/983,607
FILING DATE: April 27, 1998
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/IB96/01032
FILING DATE: June 28, 1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mary M. Krinsky
REGISTRATION NUMBER: 32423
REFERENCE/DOCKET NUMBER: OCR-679
TELECOMMUNICATION INFORMATION:
TELEPHONE: 203-773-9544
TELEFAX: 203-773-1183
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 126 residues
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE:
DESCRIPTION: polypeptide
ORIGINAL SOURCE:
ORGANISM: Homo sapiens (melanoma patient immu-
ORGANISM: nized with autologous tumor cells)
INDIVIDUAL ISOLATE: peripheral blood lymphocytes
IMMEDIATE SOURCE:
LIBRARY: DM414 scFv antibodies obtained from
LIBRARY: fuses fusion phage construct
CLONE: Z78
FEATURE:
NAME/KEY: heavy chain
US-08-983-607-26
Query Match 65.6%; Score 63; DB 3; Length 126;
Best Local Similarity 81.2%; Pred. No. 0.081;
Matches 13; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 2 INNWGGSTGYADSVKG 17
DB 51 ISNGGGSTYYADSVKG 66
RESULT 11
US-09-157-370-1
Sequence 1, Application US/09157370A
Patent No. 626238
GENERAL INFORMATION:
APPLICANT: STEINPE, Boris
APPLICANT: STEINBACHER, Stefan
TITLE OF INVENTION: PROCESS FOR MODIFYING THE STABILITY OF ANTIBODIES
FILE REFERENCE: P8341-8072
CURRENT APPLICATION NUMBER: US/09/157,370A
CURRENT FILING DATE: 1998-09-21
EARLIER APPLICATION NUMBER: 08/765,179
EARLIER FILING DATE: 1997-01-14
EARLIER APPLICATION NUMBER: PCT/EP95/02626
EARLIER FILING DATE: 1995-07-06

EARLIER APPLICATION NUMBER: DE/P44 25 115.7
EARLIER FILING DATE: 1994-07-15
NUMBER OF SEQ ID NOS: 10
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 1
LENGTH: 117
TYPE: PRT
ORGANISM: Homo sapiens
US-09-157-370-1
Query Match 65.1%; Score 62.5; DB 3; Length 117;
Best Local Similarity 59.1%; Pred. No. 0.088;
Matches 13; Conservative 1; Mismatches 3; Indels 5; Gaps 1;
QY 1 GINW-----NGGSTGYADSVKG 17
DB 44 GLEWVGWIYNGGDTYYADSVKG 65
RESULT 12
US-08-264-093-22
Sequence 22, Application US/08264093
Patent No. 5639863
GENERAL INFORMATION:
APPLICANT: Michael D. Dan
TITLE OF INVENTION: HUMAN MONOCLONAL ANTIBODIES SPECIFIC TO
TITLE OF INVENTION: CELL CYCLE-INDEPENDENT GLIOMA SURFACE
TITLE OF INVENTION: ANTIGEN
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:
ADDRESSEE: Rigout & Maybee
STREET: 2300 Richmond-Adelaide Centre
STREET: 101 Richmond Street West
CITY: Toronto
STATE: Ontario
COUNTRY: Canada
ZIP: M5H 2J7
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette - 3.5 inch, 1.4 Mb storage
COMPUTER: IBM PC Compatible
OPERATING SYSTEM: MS-DOS 6.00
SOFTWARE: ASCII Editor
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/264,093
FILING DATE:
CLASSIFICATION: 536
PRIOR APPLICATION DATA: No. 5639863 applicable
ATTORNEY/AGENT INFORMATION:
NAME: Lake, James R.
REGISTRATION NUMBER: 31081
REFERENCE/DOCKET NUMBER: NOVOP/106A/7551
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 868-1482
TELEFAX: (416) 362-0823
INFORMATION FOR SEQ ID NO: 22:
SEQUENCE CHARACTERISTICS:
LENGTH: 17 amino acids
TYPE: amino acid
STRANDEDNESS: not applicable
TOPOLOGY: linear
US-08-264-093-22
Query Match 64.8%; Score 62; DB 1; Length 17;
Best Local Similarity 81.2%; Pred. No. 0.014;
Matches 13; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 2 INNWGGSTGYADSVKG 17
DB 2 ISSNGGSTYYADSVKG 17
RESULT 13
US-09-899-896-7

; Sequence 7, Application US/09899896
; Patent No. 6569431
; GENERAL INFORMATION:
; APPLICANT: von Edingen, Hans-Christian
; APPLICANT: Genain, Claude P.
; APPLICANT: Hauser, Stephen L.
; TITLE OF INVENTION: Recombinant Antibody Fragments as Autoantibody
; TITLE OF INVENTION: Antagonists
; FILE REFERENCE: SF01-025-2
; CURRENT APPLICATION NUMBER: US/09/899,896
; CURRENT FILING DATE: 2001-07-05
; PRIOR APPLICATION NUMBER: 09/691,654
; PRIOR FILING DATE: 2000-10-17
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 7
; LENGTH: 111
; TYPE: PRT
; ORGANISM: human
US-09-899-896-7

Query Match 64.6%; Score 62; DB 4; Length 111;
Best Local Similarity 75.0%; Pred. No. 0.098;
Matches 12; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 INWNGSTGYADSVKG 17
|:::|||||
DB 51 ISYDGGSTYADSVKG 66

RESULT 14
US-08-652-816A-11
; Sequence 11, Application US/08652816A
; Patent No. 5872215
; GENERAL INFORMATION:
; APPLICANT: Osbourn, JK
; APPLICANT: Allen, DJ
; APPLICANT: McCafferty, JG
; TITLE OF INVENTION: Specific binding members, materials and
; TITLE OF INVENTION: methods.
; NUMBER OF SEQUENCES: 53
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,816A
; FILING DATE: 23-MAY-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9125579.4
; FILING DATE: 02-DEC-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9125579.8
; FILING DATE: 02-DEC-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9206318.9
; FILING DATE: 24-MAR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9206372.6
; FILING DATE: 23-SEP-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9525004.9
; FILING DATE: 07-DEC-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9610824.6
; FILING DATE: 23-MAY-1996

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB92/02240
; FILING DATE: 02-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/244,597
; FILING DATE: 01-JUN-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: David W. Clough
; REGISTRATION NUMBER: 36,107
; REFERENCE/DOCKET NUMBER: 28111/33308
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 312-474-6300
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 118 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
US-08-652-816A-11

Query Match 64.6%; Score 62; DB 2; Length 118;
Best Local Similarity 81.2%; Pred. No. 0.11;
Matches 13; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 INWNGSTGYADSVKG 17
|:|||||
DB 51 ISSNGSTYADSVKG 66

RESULT 15
US-08-264-093-14
; Sequence 14, Application US/08264093
; Patent No. 5639863
; GENERAL INFORMATION:
; APPLICANT: Michael D. Dan
; TITLE OF INVENTION: HUMAN MONOCLONAL ANTIBODIES SPECIFIC TO
; TITLE OF INVENTION: CELL CYCLE-INDEPENDENT GLIOMA SURFACE
; NUMBER OF SEQUENCES: 26
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ridout & Maybee
; STREET: 2300 Richmond-Adelaide Centre
; STREET: 101 Richmond Street West
; CITY: Toronto
; STATE: Ontario
; COUNTRY: Canada
; ZIP: M5H 2J7
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette - 3.5 inch, 1.4 Mb storage
; COMPUTER: IBM PC Compatible
; OPERATING SYSTEM: MS-DOS 6.00
; SOFTWARE: ASCII Editor
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/264,093
; FILING DATE:
; CLASSIFICATION: 536
; PRIOR APPLICATION DATA: No. 5639863 applicable
; ATTORNEY/AGENT INFORMATION:
; NAME: Lake, James R.
; REGISTRATION NUMBER: 31081
; REFERENCE/DOCKET NUMBER: NOVOP/106A/7551
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (416) 868-1482
; TELEFAX: (416) 362-0823
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 120 amino acids
; TYPE: amino acid
; STRANDEDNESS: not applicable
; TOPOLOGY: linear
US-08-264-093-14
Query Match 64.6%; Score 62; DB 1; Length 120;
Best Local Similarity 81.2%; Pred. No. 0.11;

Matches 13; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 INWNGGSGTYADSVKG 17

Db 44 ISSNGGSGTYADSVKG 59

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Job time : 40.4643 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 18, 2004, 03:46:39 ; Search time 675.143 Seconds
(without alignments)
8.086 Million cell updates/sec

Title: US-10-029-926B-115

Perfect score: 96

Sequence: 1 GINWNGSGTGADSVK 17

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Gapop 10.0 , Gapext 0.5

Searched: 1342398 seqs, 321133274 residues

Total number of hits satisfying chosen parameters: 1342398

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Published Applications AA:*

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13: /cgn2_6/ptodata/2/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/2/pubpaa/US10B_PUBCOMB.pep.*
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17: /cgn2_6/ptodata/2/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	96	100.0	17	15	US-10-032-037B-115
3	96	100.0	17	15	US-10-029-988B-115
4	96	100.0	17	15	US-10-032-423A-115
5	96	100.0	98	12	US-10-453-698-60
6	96	100.0	98	12	US-10-029-926B-61
7	96	100.0	98	14	US-10-194-975-20
8	96	100.0	98	15	US-10-308-817-60
9	96	100.0	98	15	US-10-032-037B-61
10	96	100.0	98	15	US-10-029-988B-61
11	96	100.0	98	15	US-10-032-423A-61
12	96	100.0	98	16	US-10-379-392-21
13	96	100.0	122	15	US-10-447-331-6
14	96	100.0	239	10	US-09-880-748-937
15	96	100.0	239	10	US-09-880-748-2015

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16 100.0 239 10 US-09-880-748-2038 Sequence 2038, Ap
17 100.0 239 12 US-10-293-418-937 Sequence 937, Ap
18 100.0 239 12 US-10-293-418-2015 Sequence 2015, Ap
19 100.0 239 12 US-10-293-418-2038 Sequence 2038, Ap
20 100.0 244 14 US-10-322-673-42 Sequence 42, Appl
21 100.0 286 15 US-10-032-037B-204 Sequence 204, App
22 100.0 266 15 US-10-029-988B-204 Sequence 204, App
23 100.0 266 15 US-10-032-423A-204 Sequence 204, App
24 100.0 277 12 US-10-029-926B-25 Sequence 25, Appl
25 100.0 277 12 US-10-029-926B-203 Sequence 203, App
26 100.0 277 15 US-10-032-037B-25 Sequence 25, Appl
27 100.0 277 15 US-10-032-037B-203 Sequence 203, App
28 100.0 277 15 US-10-029-988B-25 Sequence 25, Appl
29 100.0 277 15 US-10-029-988B-203 Sequence 203, App
30 100.0 277 15 US-10-032-423A-25 Sequence 25, Appl
31 100.0 277 15 US-10-032-423A-203 Sequence 203, App
32 100.0 309 13 US-10-052-798-9 Sequence 9, Appl
33 100.0 309 14 US-10-288-917-9 Sequence 9, Appl
34 100.0 309 15 US-10-423-448-9 Sequence 26, Appl
35 100.0 464 12 US-10-029-926B-26 Sequence 26, Appl
36 100.0 464 15 US-10-032-037B-26 Sequence 26, Appl
37 100.0 464 15 US-10-029-988B-26 Sequence 26, Appl
38 100.0 464 15 US-10-032-423A-26 Sequence 26, Appl
39 82.3 98 12 US-10-029-926B-60 Sequence 60, Appl
40 82.3 98 15 US-10-032-037B-60 Sequence 60, Appl
41 82.3 98 15 US-10-029-988B-60 Sequence 60, Appl
42 82.3 98 15 US-10-032-423A-60 Sequence 60, Appl
43 82.3 99 12 US-10-453-698-56 Sequence 56, Appl
44 82.3 99 14 US-10-194-975-16 Sequence 16, Appl
45 82.3 99 15 US-10-308-817-56 Sequence 56, Appl

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ALIGNMENTS

RESULT 1

US-10-029-926B-115
; Sequence 115, Application US/10029926B
; Publication No. US20040073011A1

; GENERAL INFORMATION:
; APPLICANT: HAGAY, et al.

; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
; FILE REFERENCE: 10793/50
; CURRENT APPLICATION NUMBER: US/10/029,926B

; CURRENT FILING DATE: 2001-12-31

; PRIOR APPLICATION NUMBER: 60/258,948

; PRIOR FILING DATE: 12/29/2000

; NUMBER OF SEQ ID NOS: 203

; SOFTWARE: FastSeq for Windows Version 3.0

; SEQ ID NO 115

; LENGTH: 17

; TYPE: PRT

; ORGANISM: Homo sapiens

US-10-029-926B-115

Query Match 100.0%; Score 96; DB 12; Length 17;
Best Local Similarity 100.0%; Pred. No. 4.3e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVK 17

Db 1 GINWNGSGTGADSVK 17

RESULT 2

US-10-032-037B-115
; Sequence 115, Application US/10032037B
; Publication No. US20040001822A1

; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.

; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED

; MOIETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/44

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; CURRENT APPLICATION NUMBER: US/10/032,037B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 115
; LENGTH: 17
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-032-037B-115

Query Match      100.0%; Score 96; DB 15; Length 17;
Best Local Similarity 100.0%; Pred. No. 4.3e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
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Db 1 GINWNGSGTGADSVKG 17
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RESULT 3
US-10-029-988B-115
; Sequence 115, Application US/10029988B
; Publication No. US20040001839A1
; GENERAL INFORMATION:
; APPLICANT: Bio-Technology General Corp.
; TITLE OF INVENTION: Y17-ISOLATED MOLECULES COMPRISING EPITOPES CONTAINING SULFATED
; TITLE OF INVENTION: MOIETIES, ANTIBODIES TO SUCH EPITOPES, AND USES THEREOF
; FILE REFERENCE: 10793/46
; CURRENT APPLICATION NUMBER: US/10/029,988B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 2000-12-29
; NUMBER OF SEQ ID NOS: 204
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 115
; LENGTH: 17
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-988B-115

Query Match      100.0%; Score 96; DB 15; Length 17;
Best Local Similarity 100.0%; Pred. No. 4.3e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
   |||||
Db 1 GINWNGSGTGADSVKG 17
   |||||

US-10-029-926b-115.rapb
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Best Local Similarity 100.0%; Pred. No. 4.3e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
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Db 1 GINWNGSGTGADSVKG 17
   |||||

RESULT 5
US-10-453-698-60
; Sequence 60, Application US/10453698
; Publication No. US20040038308A1
; GENERAL INFORMATION:
; APPLICANT: Rother, Russell
; TITLE OF INVENTION: HYBRID ANTIBODIES
; FILE REFERENCE: 82 CIP (1087-37 CIP)
; CURRENT APPLICATION NUMBER: US/10/453,698
; CURRENT FILING DATE: 2003-06-03
; NUMBER OF SEQ ID NOS: 196
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 60
; LENGTH: 98
; TYPE: PRT
; ORGANISM: human
US-10-453-698-60

Query Match      100.0%; Score 96; DB 12; Length 98;
Best Local Similarity 100.0%; Pred. No. 2.5e-06;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
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Db 50 GINWNGSGTGADSVKG 66
   |||||

RESULT 6
US-10-029-926B-61
; Sequence 61, Application US/10029926B
; Publication No. US20040073011A1
; GENERAL INFORMATION:
; APPLICANT: HAGAY, et al.
; TITLE OF INVENTION: SPECIFIC HUMAN ANTIBODIES FOR SELECTIVE CANCER THERAPY
; FILE REFERENCE: 10793/50
; CURRENT APPLICATION NUMBER: US/10/029,926B
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: 60/258,948
; PRIOR FILING DATE: 12/29/2000
; NUMBER OF SEQ ID NOS: 203
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 61
; LENGTH: 98
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-029-926B-61

Query Match      100.0%; Score 96; DB 12; Length 98;
Best Local Similarity 100.0%; Pred. No. 2.5e-06;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
   |||||
Db 50 GINWNGSGTGADSVKG 66
   |||||

RESULT 7
US-10-194-975-20
; Sequence 20, Application US/10194975
; Publication No. US20030039649A1
; GENERAL INFORMATION:
; APPLICANT: Foote, Jefferson
; TITLE OF INVENTION: Super Humanized Antibodies
; FILE REFERENCE: 501231.01
; CURRENT APPLICATION NUMBER: US/10/194,975
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Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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APPLICANT: Lazar, Gregory Alan
; APPLICANT: Desjarlais, John Rudolf
; APPLICANT: Marshall, Shannon Alicia

```
; APPLICANT: Dahiyat, Bassil I.
; TITLE OF INVENTION: ANTIBODY OPTIMIZATION
; FILE REFERENCE: A-71386-3 463077-236
; CURRENT APPLICATION NUMBER: US/10/379,392
; CURRENT FILING DATE: 2003-03-03
; PRIOR APPLICATION NUMBER: US 60/360,843
; PRIOR FILING DATE: 2002-03-01
; PRIOR APPLICATION NUMBER: US 60/384,197
; PRIOR FILING DATE: 2002-05-29
; NUMBER OF SEQ ID NOS: 184
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 21
; LENGTH: 98
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-379-392-21

Query Match      100.0%; Score 96; DB 16; Length 98;
Best Local Similarity 100.0%; Pred. No. 2.5e-06;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
Db 50 GINWNGSGTGADSVKG 66

RESULT 13
US-10-447-331-6
; Sequence 6, Application US/10447331
; Publication No. US20030219434A1
; GENERAL INFORMATION:
; APPLICANT: Garter, Paul J.
; APPLICANT: Ridgway, John B.
; TITLE OF INVENTION: ANTIBODIES FOR CANCER THERAPY AND
; TITLE OF INVENTION: DIAGNOSIS
; FILE REFERENCE: GENENT.122A
; CURRENT APPLICATION NUMBER: US/10/447,331
; CURRENT FILING DATE: 2003-05-28
; PRIOR APPLICATION NUMBER: US/09/515,825
; PRIOR FILING DATE: 2000-02-29
; PRIOR APPLICATION NUMBER: 60/122262
; PRIOR FILING DATE: 1999-03-01
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 122
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-447-331-6

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Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17
Db 50 GINWNGSGTGADSVKG 66

RESULT 14
US-09-880-748-937
; Sequence 937, Application US/09880748
; Publication No. US20030059937A1
; GENERAL INFORMATION:
; APPLICANT: Ruben et al.
; TITLE OF INVENTION: Antibodies that Immunospecifically Bind Blys
; FILE REFERENCE: PF523
; CURRENT APPLICATION NUMBER: US/09/880,748
; CURRENT FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/212,210
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: 60/240,816
; PRIOR FILING DATE: 2001-03-16
; PRIOR APPLICATION NUMBER: 60/277,379
; PRIOR FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: 60/293,499
; PRIOR FILING DATE: 2001-05-25
; NUMBER OF SEQ ID NOS: 3239
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2015
; LENGTH: 239
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-880-748-937

Query Match      100.0%; Score 96; DB 10; Length 239;
Best Local Similarity 100.0%; Pred. No. 6.1e-06;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 50 GINWNGSGTGADSVKG 66

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Job time : 675.143 secs
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GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 18, 2004, 03:33:04; Search time 34.6071 Seconds
(without alignments)
47.252 Million cell updates/sec

Title: US-10-029-926B-115
Perfect score: 96
Sequence: 1 GINWNGSGTGYADSVKG 17

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Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: PIR 78: *
1: pir1: *
2: pir2: *
3: pir3: *
4: pir4: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match	Score	Length	DB ID	Description
1	96	100.0	98	2 S26928	Ig heavy chain V r
2	87	90.6	112	2 PH1654	Ig heavy chain V r
3	79	82.3	98	2 S26927	Ig heavy chain V r
4	79	82.3	121	2 S31104	Ig heavy chain (su
5	79	82.3	128	2 S31595	Ig heavy chain V r
6	76	75.2	123	2 S30532	Ig heavy chain V r
7	74	77.1	121	2 S31118	Ig heavy chain - h
8	73	76.0	100	2 S69896	Ig heavy chain V r
9	71	74.0	98	2 S26929	Ig heavy chain V r
10	71	74.0	98	2 S26934	Ig heavy chain V r
11	71	74.0	120	2 S44111	Ig heavy chain V-D
12	71	74.0	191	2 JF0048	Ig heavy chain V r
13	69	71.9	120	1 GH0048	Ig heavy chain V r
14	68	70.8	145	2 S11239	Ig heavy chain V r
15	65	67.7	120	2 S36273	Ig heavy chain V r
16	65	67.7	123	2 PC4281	Ig heavy chain V r
17	64	66.7	90	2 S24248	anti-SS-A/Ro 60K p
18	64	66.7	120	2 S26278	Ig heavy chain V r
19	62	64.6	96	2 S20781	Ig heavy chain V r
20	61.5	64.1	97	2 S26935	Ig heavy chain V r
21	61.5	64.1	97	2 S46462	Ig heavy chain V r
22	61.5	64.1	116	2 B28966	Ig heavy chain V r
23	61.5	64.1	116	2 S31110	Ig heavy chain pre
24	61.5	64.1	146	2 I47184	Ig heavy chain - h
25	61	63.5	98	2 S26889	Ig heavy chain var
26	61	63.5	98	2 S24858	Ig heavy chain V r
27	61	63.5	99	2 S24259	Ig heavy chain V r
28	61	63.5	100	2 S24258	Ig heavy chain V r
29	61	63.5	101	2 S24257	Ig heavy chain V r

30 61 63.5 102 2 S24260 Ig heavy chain V r
31 61 63.5 104 2 S24255 Ig heavy chain V r
32 61 63.5 105 2 S24249 Ig heavy chain V r
33 61 63.5 105 2 S24256 Ig heavy chain V r
34 61 63.5 108 2 PH1648 Ig heavy chain V r
35 61 63.5 109 2 PH1649 Ig heavy chain V r
36 61 63.5 109 2 S24254 Ig heavy chain V r
37 61 63.5 109 2 S24253 Ig heavy chain V r
38 61 63.5 110 2 S24250 Ig heavy chain V r
39 61 63.5 112 2 PH1647 Ig heavy chain V r
40 61 63.5 113 2 S24247 Ig heavy chain V r
41 61 63.5 115 2 S20982 Ig heavy chain - c
42 61 63.5 117 2 A45953 Ig heavy chain pre
43 61 63.5 118 2 S31121 Ig heavy chain - h
44 61 63.5 119 2 C36005 Ig heavy chain V r
45 61 63.5 119 2 D36005 Ig heavy chain V r

ALIGNMENTS

RESULT 1

S26928
Ig heavy chain V region (DP-32) - human (fragment)
C/Species: Homo sapiens (man)
C/Date: 22-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 23-Jul-1999
C/Accession: S26928
R/Tomlinson, I.M.; Walter, G.; Marks, J.D.; Llewellyn, M.B.; Winter, G.
J. Mol. Biol. 227, 776-798, 1992
A/Title: The repertoire of human germline V(H) sequences reveals about fifty groups of
A/Reference number: S26885; MUID:93021117; PMID:1404398
A/Accession: S26928
A/Status: Preliminary
A/Molecule type: DNA
A/Residues: 1-98 <TOM>
A/Cross-references: EMBL:Z12334; NID:G32887; PIDN:CAA78204.1; PID:G32888
C/Superfamily: immunoglobulin V region; immunoglobulin homology
C/Keywords: heterotetramer; immunoglobulin
F:15-98/Domain: immunoglobulin homology <IMM>

Query Match 100.0%; Score 96; DB 2; Length 98;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db 50 GINWNGSGTGYADSVKG 66
|||||

RESULT 2

PH1654
Ig heavy chain V region (clone 6H9) - human (fragment)
C/Species: Homo sapiens (man)
C/Date: 24-Feb-1994 #sequence_revision 24-Feb-1994 #text_change 16-Aug-1996
C/Accession: PH1654
R/Hillson, J.L.; Karr, N.S.; Opplinger, I.R.; Mannik, M.; Sasso, E.H.
J. Exp. Med. 178, 331-336, 1993
A/Title: The structural basis of germline-encoded VH3 immunoglobulin binding to staphyl
A/Reference number: PH1642; MUID:93301610; PMID:8315388
A/Accession: PH1654
A/Molecule type: mRNA
A/Residues: 1-112 <HIL>
A/Experimental source: B cell
C/Superfamily: immunoglobulin V region; immunoglobulin homology
C/Keywords: heterotetramer; immunoglobulin
F:7-90/Domain: immunoglobulin homology <IMM>

Query Match 90.6%; Score 87; DB 2; Length 112;
Best Local Similarity 94.1%; Pred. No. 5.6e-06;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
|||||

Db 42. GINWGGSTGYADSVKG 58

RESULT 3

S26927

IG heavy chain V region (DP-31) - human (fragment)

C:Species: Homo sapiens (man)

C>Date: 22-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 23-Jul-1999

C:Accession: S26927

R:Tomlinson, J.M.; Walter, G.; Marks, J.D.; Llewellyn, M.B.; Winter, G.

J. Mol. Biol. 227, 776-798, 1992

A:Title: The repertoire of human germline V(H) sequences reveals about fifty groups of V

A:Reference number: S26885; MUID:93021117; PMID:1404388

A:Accession: S26927

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-98 <TOM>

A:Cross-references: EMBL:Z12333; NID:G32985; PIDN:CAA78203.1; PID:G32886

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

F:15-98/Domain: immunoglobulin homology <IMM>

Query Match 82.3%; Score 79; DB 2; Length 98;

Best Local Similarity 82.4%; Pred. No. 7.8e-05;

Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 GINWGGSTGYADSVKG 17

Db 50 GISWNSGIGYADSVKG 66

RESULT 4

S31104

IG heavy chain (subclass IgM) - human (fragment)

C:Species: Homo sapiens (man)

C>Date: 22-Nov-1993 #sequence_revision 26-May-1995 #text_change 23-Jul-1999

C:Accession: S31104

R:Raaphorst, F.M.; Timmers, E.; Kenter, M.J.H.; van Tol, M.J.D.; Vossen, J.M.; Schuurman

Eur. J. Immunol. 22, 247-251, 1992

A:Title: Restricted utilization of germ-line V(H)3 genes and short diverse third comple

A:Reference number: S31104; MUID:92111633; PMID:1730252

A:Accession: S31104

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: mRNA

A:Residues: 1-121 <PAA>

A:Cross-references: EMBL:X63080; NID:G32646; PIDN:CAA44802.1; PID:G32647

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, October 1991

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

F:15-98/Domain: immunoglobulin homology <IMM>

Query Match 82.3%; Score 79; DB 2; Length 121;

Best Local Similarity 82.4%; Pred. No. 9.6e-05;

Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 GINWGGSTGYADSVKG 17

Db 50 GISWNSGIGYADSVKG 66

RESULT 5

S31595

IG heavy chain V region - human

C:Species: Homo sapiens (man)

C>Date: 03-Mar-1994 #sequence_revision 10-Nov-1995 #text_change 23-Jul-1999

C:Accession: S31595

R:Cuisinier, A.M.; Gauthier, L.; Boublil, L.; Fougereau, M.; Tonnelle, C.

submitted to the EMBL Data Library, June 1992

A:Description: Mechanisms that generate human immunoglobulin diversity operate from the

A:Reference number: S31585

A:Accession: S31595

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-128 <CUI>

A:Cross-references: EMBL:Z14171; NID:G31007; PIDN:CAA78540.1; PID:G31008

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

F:23-106/Domain: immunoglobulin homology <IMM>

Query Match 82.3%; Score 79; DB 2; Length 128;

Best Local Similarity 82.4%; Pred. No. 0.0001;

Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 GINWGGSTGYADSVKG 17

Db 58 GISWNSGIGYADSVKG 74

RESULT 6

S30532

IG heavy chain V region - human

C:Species: Homo sapiens (man)

C>Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 16-Aug-1996

C:Accession: S30532

R:Mariette, X.

submitted to the EMBL Data Library, October 1992

A:Reference number: S30520

A:Accession: S30532

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-123 <MAP>

A:Cross-references: EMBL:Z18318

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

F:15-98/Domain: immunoglobulin homology <IMM>

Query Match 79.2%; Score 76; DB 2; Length 123;

Best Local Similarity 76.5%; Pred. No. 0.00027;

Matches 13; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 GINWGGSTGYADSVKG 17

Db 50 GISWNSGIGYADSVKG 66

RESULT 7

S31118

IG heavy chain - human

C:Species: Homo sapiens (man)

C>Date: 02-Dec-1993 #sequence_revision 26-May-1995 #text_change 17-Mar-1999

C:Accession: S31118

R:Raaphorst, F.M.; Timmers, E.; Kenter, M.J.H.; van Tol, M.J.D.; Vossen, J.M.; Schuurman

Eur. J. Immunol. 22, 247-251, 1992

A:Title: Restricted utilization of germ-line V(H)3 genes and short diverse third comple

A:Reference number: S31104; MUID:92111633; PMID:1730252

A:Accession: S31118

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: mRNA

A:Residues: 1-121 <PAA>

A:Cross-references: EMBL:X62969

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, October 1991

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

F:15-98/Domain: immunoglobulin homology <IMM>

Query Match 77.1%; Score 74; DB 2; Length 121;

Best Local Similarity 76.5%; Pred. No. 0.00054;

Matches 13; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 GINWGGSTGYADSVKG 17

Db 50 GISWNSGIGYADSVKG 66

RESULT 8

S69896

RECORD 13
G1HUDB

```

Ig heavy chain V-III region (Dob) - human
C:Species: Homo sapiens (man)
C:Date: 31-Dec-1979 #sequence_revision 31-Dec-1979 #text_change 02-Sep-1997
C:Accession: A90431; A02065
R:Steiner, L.A.; Garcia Pardo, A.; Margolies, M.N.
Biochemistry 18, 4068-4080, 1979
A:Title: Amino acid sequence of the heavy-chain variable region of the crystallizable hu
A:Reference number: A90431; MUID:80020921; PMID:114209
A:Accession: A90431
A:Molecule type: protein
A:Residues: 1-120 <STE>
R:Steiner, L.A.; Lopes, A.D.
Biochemistry 18, 4054-4067, 1979
A:Title: The crystallizable human myeloma protein Dob has a hinge-region deletion.
A:Reference number: A90430; MUID:80020920; PMID:114208
A:Contents: annotation
A:Note: this gamma-1 myeloma protein has a deletion in the hinge region; there are no li
C:Genetics:
A:Gene: GDB:IGHV6
A:Cross-references: GDB:128528; OMIM:147070
A:Map position: 14q32.33-14q32.33
C:Superfamily: immunoglobulin V region; immunoglobulin homology
C:Keywords: heterotetramer; immunoglobulin
F:15-98/Domain: immunoglobulin homology <IMM>
F:22-96/Disulfide bonds: #status predicted

Query Match 71.9%; Score 69; DB 1; Length 120;
Best Local Similarity 81.2%; Pred. No. 0.003; 3; Indels 0; Gaps 0;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 INWNGSGTGYADSVKG 17
Db 51 ITWNGSVLYADSVKG 66

RESULT 14
S11239
IG heavy chain V region - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 21-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 23-Jul-1999
C:Accession: S11239
R:Feigenhauer, M.; Kohl, J.; Rueker, F.
Nucleic Acids Res. 18, 4927, 1990
A:Title: Nucleotide sequences of the cDNAs encoding the V-regions of H- and L-chains of
A:Reference number: S11239; MUID:90370490; PMID:1637678
A:Accession: S11239
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-145 <FEL>
A:Cross-references: EMBL:X53613; NID:G23865; PIDN:CAA37675.1; PID:G762936
C:Superfamily: immunoglobulin V region; immunoglobulin homology
C:Keywords: heterotetramer; immunoglobulin
F:34-117/Domain: immunoglobulin homology <IMM>

Query Match 70.8%; Score 68; DB 2; Length 145;
Best Local Similarity 70.6%; Pred. No. 0.0051; 3; Indels 0; Gaps 0;
Matches 12; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db 69 GISWDSSSIGYADSVKG 85

RESULT 15
S36273
IG heavy chain V region (clone alpha-THY-32) - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 23-Jul-1999
C:Accession: S36273
R:Griffiths, A.D.; Malmqvist, M.; Marks, J.D.; Bye, J.M.; Embleton, M.J.; McCafferty, J.
EMBO J. 12, 723-734, 1993
A:Title: Human anti-self antibodies with high specificity from phage display libraries.
A:Reference number: S36256; MUID:93178448; PMID:7679990

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A:Accession: S36273
A>Status: preliminary; nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-120 <GRI>
A:Cross-references: EMBL:Z18834; NID:G33116; PIDN:CAA79286.1; PID:G939896
C:Superfamily: immunoglobulin V region; immunoglobulin homology
C:Keywords: heterotetramer; immunoglobulin
F:15-98/Domain: immunoglobulin homology <IMM>

Query Match 67.7%; Score 65; DB 2; Length 120;
Best Local Similarity 76.5%; Pred. No. 0.012;
Matches 13; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 GINWNGSGTGYADSVKG 17
Db 50 GISSNSGSIGYADSVKG 66

Search completed: September 18, 2004, 03:47:33
Job time : 35.6071 secs

```


GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 18, 2004, 02:29:47 ; Search time 24.8929 Seconds
(without alignments)
35.560 Million cell updates/sec

Title: US-10-029-926B-115

Perfect score: 96

Sequence: 1 GINWGGSTGYADSVK 17

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	69	71.9	120	1 HV3U_HUMAN	P01782 homo sapien
2	61.5	64.1	116	1 HV05_CARAU	P19181 carassius a
3	59	61.5	117	1 HV53_MOUSE	P18524 mus musculu
4	57	59.4	117	1 HV3C_HUMAN	P01764 homo sapien
5	53	55.2	115	1 HV3F_HUMAN	P01767 homo sapien
6	51	53.1	121	1 HV3J_HUMAN	P01771 homo sapien
7	50.5	52.6	97	1 HV56_MOUSE	P18527 mus musculu
8	48.5	50.5	117	1 HV02_CANFA	P01785 mus musculu
9	48	50.0	117	1 HV55_MOUSE	P18526 mus musculu
10	48	50.0	119	1 HV3L_HUMAN	P01773 homo sapien
11	47	49.0	117	1 HV54_MOUSE	P18525 mus musculu
12	46.5	48.4	135	1 HV02_XENLA	P20957 xenopus lae
13	46	47.9	114	1 HV3E_HUMAN	P01783 homo sapien
14	46	47.9	115	1 HV3D_HUMAN	P01785 homo sapien
15	46	47.9	117	1 HV03_CARAU	P19180 carassius a
16	46	47.9	119	1 HV3M_HUMAN	P01774 homo sapien
17	46	47.9	119	1 HV3N_HUMAN	P01775 homo sapien
18	46	47.9	122	1 HV3A_HUMAN	P01762 homo sapien
19	46	47.9	126	1 HV3K_HUMAN	P01772 homo sapien
20	46	47.9	428	1 SP7_MOUSE	Q8vi67 mus musculu
21	46	47.9	431	1 SP7_HUMAN	Q8td32 homo sapien
22	46	47.9	559	1 PMGI_MAIZE	P30792 zea mays (m
23	45.5	47.4	111	1 HV35_MOUSE	P01804 mus musculu
24	45	46.9	116	1 HV3T_HUMAN	P01781 homo sapien
25	45	46.9	116	1 HV45_MOUSE	P01821 mus musculu
26	45	46.9	117	1 HV1B_HUMAN	P01743 homo sapien
27	45	46.9	121	1 HV01_MOUSE	P01745 mus musculu
28	45	46.9	122	1 HV3G_HUMAN	P01768 homo sapien
29	45	46.9	954	1 XYNA_RUMFL	P29126 ruminococcu
30	44.5	46.4	120	1 HV3E_HUMAN	P01766 homo sapien
31	44	45.8	115	1 HV32_MOUSE	P01801 mus musculu
32	44	45.8	725	1 AREA_PENCH	Q01582 penicillium
33	44	45.8	865	1 NRFA_PENUR	Q92269 penicillium

34	43	44.8	136	1 HV16_MOUSE	P01783 mus musculu
35	43	44.8	151	1 YF84_METUA	Q58979 methanococc
36	43	44.8	327	1 K6P1_BACTN	Q848r5 bacteroides
37	43	44.8	527	1 PPCK_OCEIH	Q84p04 oceanobacil
38	43	44.8	549	1 PRIS_VACMU	Q28520 macaca mula
39	43	44.8	676	1 PRIS_HUMAN	P07225 homo sapien
40	43	44.8	678	1 GSPD_AERSA	P45778 aeromonas s
41	43	44.8	731	1 HUTU_CABEL	Q9nae2 caenothabdi
42	43	44.8	835	1 FASD_ECOLI	P46000 escherichia
43	43	44.8	1019	1 CAL6_CHICK	P20785 gallus gall
44	43	44.8	1633	1 YP74_CABEL	Q09221 caenothabdi
45	42.5	44.3	264	1 EXB5_ARATH	Q9m203 arabidopsi

ALIGNMENTS

RESULT 1					
HV3U_HUMAN					
ID	HV3U_HUMAN	STANDARD;	PRT;	120 AA.	
AC	P01782;				
DT	21-JUL-1986 (Rel. 01, Created)				
DT	21-JUL-1986 (Rel. 01, Last sequence update)				
DT	10-OCT-2003 (Rel. 42, Last annotation update)				
DE	Ig heavy chain V-III region DOB.				
OS	Homo sapiens (Human)				
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
OX	NCBI_TaxID=9606;				
RN	[1]				
RP	SEQUENCE.				
RX	MEDLINE=80020921; PubMed=114209;				
RA	Steiner L.A., Garcia Pardo A., Margolies M.N.;				
RT	"Amino acid sequence of the heavy-chain variable region of the				
RT	crystallizable human myeloma protein Dob.";				
RL	Biochemistry 18:4068-4080(1979).				
RN	[2]				
RP	CRYSTALLIZATION.				
RX	MEDLINE=80020920; PubMed=114208;				
RA	Steiner L.A., Lopes A.D.;				
RT	"The crystallizable human myeloma protein Dob has a hinge-region				
RT	deletion.";				
CC	-1- MISCELLANEOUS: THIS GAMMA-1 MYELOMA PROTEIN HAS A DELETION IN THE				
CC	HINGE REGION. THERE ARE NO LIGHT-HEAVY OR INTER-HEAVY CHAIN				
CC	DISULFIDE BONDS.				
CC	-1- SIMILARITY: Contains 1 immunoglobulin-like domain.				
DR	PIR; A90431; GHUDB.				
DR	HSSP; P01772; 2FB4.				
DR	GO; GO:0005576; C:extracellular; NAS.				
DR	GO; GO:0003823; F:antigen binding; NAS.				
DR	GO; GO:0006955; P:immune response; NAS.				
DR	InterPro; IPR007110; IG-like.				
DR	InterPro; IPR003596; IG_V.				
DR	SMART; SM00406; IGV; 1.				
DR	PROSITE; PS00835; IG LIKE; 1.				
DR	Immunoglobulin V region.				
FT	DOMAIN 1 112				
FT	NON_TER 120 120				
SQ	SEQUENCE 120 AA; 13440 MW; 880DDE307C4B2627 CRC64;				

Query Match 71.9%; Score 69; DB 1; Length 120;
Best Local Similarity 81.2%; Pred. No. 0.00061;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY	2	INWNGSGTGYADSVK 17
DB	51	ITWNGGSLYADSVK 66

RESULT 2
HV05_CARAU

CC -I- MISCELLANEOUS: THIS CHAIN WAS ISOLATED FROM AN IGG1 MYELOMA

RESULT 11
HV54 MOUSE
ID HV54 MOUSE
AC P18525; PRT; 117 AA.

DT 01-NOV-1990 (Rel. 16, Last created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Ig heavy chain V region 5-84 precursor.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
CX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BALB/CJ;
RX MEDLINE=89279149; PubMed=2499654;
RA Levy N.S., Malipiero U.V., Lebecque S.G., Gearhart P.J.;
RT "Early onset of somatic mutation in immunoglobulin VH genes during
RT the primary immune response.";
RL J. Exp. Med. 169:2007-2019(1989).
CC -1- MISCELLANEOUS: THIS SEQUENCE BELONGS TO THE VH7183 SUBFAMILY.
DR PIR; J03005; HYMS84.
DR HSSP; P01810; 2FEJ.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003596; Ig_V.
DR Pfam; PF00047; Ig; 1.
DR SMART; SMC0406; IGV; 1.
DR PROSITE; PS50835; IG-LIKE; 1.
KW Immunoglobulin V region; Signal.
FT SIGNAL 19
FT CHAIN 20 117 IG HEAVY CHAIN V REGION 5-84.
FT DOMAIN 20 49 FRAMEWORK-1.
FT DOMAIN 50 54 COMPLEMENTARITY-DETERMINING-1.
FT DOMAIN 55 68 FRAMEWORK-2.
FT DOMAIN 69 85 COMPLEMENTARITY-DETERMINING-2.
FT DOMAIN 86 117 FRAMEWORK-3.
FT DISULFID 41 115 BY SIMILARITY.
FT NON_TER 117 117
SQ SEQUENCE 117 AA; 12872 MW; 234055CB6A469861 CRC64;

Query Match 49.08; Score 47; DB 1; Length 117;
Best Local Similarity 62.58; Pred. No. 1.7; Indels 0; Gaps 0;
Matches 10; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 2 INWNGSTGYADSVKG 17
Db 70 ISNGGGSTYTPDTVKG 85

RESULT 12
ID HV02 XENLA STANDARD; PRT; 135 AA.
AC P20957;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Ig heavy chain V region XIG14 precursor (Fragment).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=88176921; PubMed=2451244;
RA Schwager J., Mikoryak C.A., Steiner L.A.;
RT "Amino acid sequence of heavy chain from Xenopus laevis IGM deduced
RT from cDNA sequence: implications for evolution of immunoglobulin
RT domains.";
RL Proc. Natl. Acad. Sci. U.S.A. 85:2245-2249(1988).
CC -1- SIMILARITY: Contains 1 immunoglobulin-like domain.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; J03632; AAA49791.1; -.
DR PIR; B31933; B31933.
DR HSSP; P01810; 2FEJ.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003596; Ig_V.
DR Pfam; PF00047; Ig; 1.
DR SMART; SMC0406; IGV; 1.
DR PROSITE; PS50835; IG-LIKE; 1.
KW Immunoglobulin V region; Signal.
FT NON_TER 1
FT SIGNAL <1 18
FT CHAIN 19 135 IG HEAVY CHAIN V REGION XIG14.
FT DOMAIN 20 128 IG-LIKE.
FT NON_TER 135 135
SQ SEQUENCE 135 AA; 15080 MW; EBC467105C00732E CRC64;

Query Match 48.44; Score 46.5; DB 1; Length 135;
Best Local Similarity 47.64; Pred. No. 2.3;
Matches 10; Conservative 2; Mismatches 4; Indels 5; Gaps 1;

Qy 1 GINW-----NGSGTGYADSVK 16
Db 61 GLEWIGVIATGSGTAIDSLK 81

RESULT 13
ID HV3B HUMAN STANDARD; PRT; 114 AA.
AC P01763;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Ig heavy chain V-III region WEA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE.
RX MEDLINE=83273707; PubMed=6410398;
RA Goni F., Frangione B.;
RT "Amino acid sequence of the Fv region of a human monoclonal Igm
RT (protein WEA) with antibody activity against 3,4-pyruvylated
RT galactose in Klebsiella polysaccharides K30 and K33.";
RL Proc. Natl. Acad. Sci. U.S.A. 80:4837-4841(1983).
CC -1- MISCELLANEOUS: THIS CHAIN WAS OBTAINED FROM A MONOCLONAL ANTIBODY
CC AGAINST 3,4-PYRUVYLATED GALACTOSE AND ISOLATED FROM A PATIENT WITH
CC WALDENSTROM'S MACROGLOBULINEMIA.
CC -1- SIMILARITY: Contains 1 immunoglobulin-like domain.
DR PIR; A02046; M3HWE.
DR HSSP; P01772; 2FB4.
DR GO; GO:0005576; C:extracellular; NAS.
DR GO; GO:0003823; F:antigen binding; NAS.
DR GO; GO:0006955; P:immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003596; Ig_V.
DR Pfam; PF00047; Ig; 1.
DR SMART; SMC0406; IGV; 1.
DR PROSITE; PS50835; IG-LIKE; 1.
KW Immunoglobulin V region; Pyroglutamate carboxylic acid.
FT DOMAIN 1 112 IG-LIKE.
FT MCD_RES 1 1
FT NON_TER 114 114 PYRROLIDONE CARBOXYLIC ACID.
SQ SEQUENCE 114 AA; 12256 MW; D88294FB18A07B7 CRC64;

Query Match 47.98; Score 46; DB 1; Length 114;
Best Local Similarity 43.58; Pred. No. 2.3;
Matches 10; Conservative 3; Mismatches 4; Indels 6; Gaps 1;

Qy 1 GINW-----NGSGTGYADSVK 17

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OM protein - protein search, using sw model

Run on: September 18, 2004, 03:24:30 ; Search time 88.6429 Seconds
(without alignments)
60.510 Million cell updates/sec

Title: US-10-029-926b-115

Perfect score: 96

Sequence: 1 GINWNGSGTGADSVKG 17

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phage:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_virus:*
- 16: sp_bacteriap:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	96	100.0	112	4 Q9HCC1	Q9hcc1 homo sapien
2	84	87.5	499	4 Q8N5K4	Q8n5k4 homo sapien
3	79	82.3	104	4 Q8UL87	Q8ul87 homo sapien
4	79	82.3	573	4 Q8WU38	Q8wu38 homo sapien
5	61	63.5	121	4 Q9UL71	Q9ul71 homo sapien
6	61	63.5	597	4 Q96BB9	Q96bb9 homo sapien
7	56	58.3	118	4 Q9UL72	Q9ul72 homo sapien
8	55	57.3	479	11 Q91WP5	Q91wp5 mus musculus
9	54	56.2	112	4 Q9UGP3	Q9ugp3 homo sapien
10	51	53.1	469	2 Q9FD12	Q9fd12 shewanella
11	51	53.1	838	16 Q7UUV9	Q7uuv9 rhodospirillum rubrum
12	49	51.0	767	5 Q9U2J4	Q9u2j4 caenorhabditis elegans
13	48	50.0	279	16 Q9WZJ8	Q9wzj8 thermotoga
14	48	50.0	298	10 Q9XGB4	Q9xgb4 trifolium
15	48	50.0	416	4 Q9NP66	Q9np66 homo sapien
16	48	50.0	469	16 Q8PDZ8	Q8pdz8 xanthomonas

17	48	50.0	473	11	Q91Z05	Q91z05 mus musculus
18	48	50.0	654	5	Q86AX3	Q86ax3 dictyostell
19	48	50.0	761	5	Q20898	Q20898 caenorhabdi
20	48	50.0	4342	16	Q91157	Q91157 pseudomonas
21	47	49.0	87	9	Q7Y3K4	Q7y3k4 enterobacte
22	47	49.0	116	4	Q9UL93	Q9ul93 homo sapien
23	47	49.0	469	16	Q87CN9	Q87cn9 xylella fas
24	47	49.0	515	5	Q16968	Q16968 aplysia cal
25	47	49.0	613	4	Q8WUK1	Q8wuk1 homo sapien
26	46	47.9	113	4	Q9UL90	Q9ul90 homo sapien
27	46	47.9	122	4	Q9UL84	Q9ul84 homo sapien
28	46	47.9	131	4	Q9UL88	Q9ul88 homo sapien
29	46	47.9	159	16	Q92CR6	Q92cr6 listeria in
30	46	47.9	159	16	Q8Y7X8	Q8y7x8 listeria in
31	46	47.9	260	16	Q8NPS3	Q8nps3 corynebacte
32	46	47.9	413	4	Q7Z718	Q7z718 homo sapien
33	46	47.9	428	11	Q811U1	Q811u1 rattus norv
34	46	47.9	464	10	Q887G0	Q887g0 oryza sativ
35	46	47.9	469	16	Q8FQW5	Q8fqw5 xanthomonas
36	46	47.9	473	2	Q9RBK1	Q9rbk1 xanthomonas
37	46	47.9	493	4	Q8NCL6	Q8nc16 homo sapien
38	46	47.9	627	16	Q8P8D7	Q8p8d7 xanthomonas
39	46	47.9	660	16	Q8PJW5	Q8pjw5 xanthomonas
40	45	47.4	666	5	Q9VA77	Q9va77 drosophila
41	45	46.9	124	4	Q9UL92	Q9ul92 homo sapien
42	45	46.9	125	4	Q9UL95	Q9ul95 homo sapien
43	45	46.9	308	2	Q9S0K5	Q9s0k5 shewanella
44	45	46.9	376	15	Q7ZP49	Q7zpz49 human immu
45	45	46.9	393	16	Q8YSL5	Q8ysl5 anabaena sp

ALIGNMENTS

RESULT 1

Q9HCC1 PRELIMINARY; PRT; 112 AA.

ID Q9HCC1
AC Q9HCC1
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DE Single chain Fv (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kikuchi M., Takeda C., Tsujimoto Y., Asada S., Nagata K.;
RT "An antibody fragment2A3 specific for native lysozyme : Isolation from a
human synthetic phage display library and characterization.";
RL Submitted (CC-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB049915; BAB16829.1; -
DR HSSP; P01772; 2FB4.
DR InterPro; IPR007110; IG-like.
DR InterPro; IPR003596; IG_v.
DR Pfam; PF00047; IG_1.
DR SMART; SM00406; IGV; 1.
DR PROSITE; PS50835; IG_LIKE; 1.
FT NON_TER 1
FT NON_TER 112
SQ SEQUENCE 112 AA; 12243 MW; 24FlA45EC3B94788 CRC64;

Query Match 100.0%; Score 96; DB 4; Length 112;
Best Local Similarity 100.0%; Pred. No. 6e-07;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GINWNGSGTGADSVKG 17

Db 50 GINWNGSGTGADSVKG 66

RESULT 2


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DR PROSITE; PS50835; IG LIKE; 1.
FT NON_TER 1
FT NON_TER 121
SQ SEQUENCE 121 AA; 13154 MW; 2F045CCFA5D50736 CRC64;

Query Match 63.5%; Score 61; DB 4; Length 121;
Best Local Similarity 56.5%; Pred. No. 0.14;
Matches 13; Conservative 2; Mismatches 6; Gaps 1;

QY 1 GINW-----NGSGTGYADSVKG 17
| : : : : :
| : : : : :
| : : : : :
44 GLEWVSLISGSGSTYYADSVKG 66

Db
RESULT 6
Q96BB9 PRELIMINARY; PRT; 597 AA.
ID Q96BB9
AC Q96BB9;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Hypothetical protein.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-B-cell;
RA Strausberg R.;
RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC015760; AAH15760.1; -.
DR InterPro; IPR007110; IG LIKE.
DR InterPro; IPR003006; IG MHC.
DR InterPro; IPR003596; IG_V.
DR Pfam; PF00047; IG; 5.
DR SMART; SM00406; IGV; 1.
DR PROSITE; PS50835; IG LIKE; 5.
DR PROSITE; PS00290; IG MHC; 3.
KW Hypothetical protein.
SQ SEQUENCE 597 AA; 65039 MW; 4FCA3AD8ECE263D9 CRC64;

Query Match 63.5%; Score 61; DB 4; Length 597;
Best Local Similarity 56.5%; Pred. No. 0.86;
Matches 13; Conservative 2; Mismatches 6; Gaps 1;

QY 1 GINW-----NGSGTGYADSVKG 17
| : : : : :
| : : : : :
| : : : : :
63 GLEWVSAISGSGSTYYADSVKG 85

Db
RESULT 7
Q9UL72 PRELIMINARY; PRT; 118 AA.
ID Q9UL72
AC Q9UL72;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Myosin-reactive immunoglobulin heavy chain variable region
DE (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98277139; PubMed=9614934;
RA You X., Liu B., Van der Merwe P.L., Kallis N.N., Berney S.M.,
RA Young D.C.;
RT "Myosin-reactive autoantibodies in rheumatic carditis and normal
RT fetus.";
RL Clin. Immunol. Immunopathol. 87:184-192(1998).
DR EMBL; AF035042; AAD56278.1; -.

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DR PIR; S21205; S21205.
DR HSSP; P01772; 2FB4.
DR InterPro; IPR007110; IG-like.
DR InterPro; IPR003596; IG_V.
DR Pfam; PF00047; IG; 1.
DR SMART; SM00406; IGV; 1.
DR PROSITE; PS50835; IG LIKE; 1.
FT NON_TER 1
FT NON_TER 118
SQ SEQUENCE 118 AA; 12872 MW; B4D1A5944B2D5CCA CRC64;

Query Match 58.3%; Score 56; DB 4; Length 118;
Best Local Similarity 62.5%; Pred. No. 0.78;
Matches 10; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 INWNGSGTGYADSVKG 17
: : : : :
: : : : :
: : : : :
50 VTYSGGSYADSVKG 65

Db
RESULT 8
Q91WP5 PRELIMINARY; PRT; 479 AA.
ID Q91WP5
AC Q91WP5;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Hypothetical protein.
DE Hypothetical protein.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Colon;
RA Strausberg R.;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC013656; AAH13656.1; -.
DR InterPro; IPR007110; IG-like.
DR InterPro; IPR003006; IG MHC.
DR InterPro; IPR003596; IG_V.
DR Pfam; PF00047; IG; 4.
DR SMART; SM00406; IGV; 1.
DR PROSITE; PS50835; IG LIKE; 4.
DR PROSITE; PS00290; IG MHC; 2.
KW Hypothetical protein.
SQ SEQUENCE 479 AA; 51603 MW; ECB2D087748584F CRC64;

Query Match 57.3%; Score 55; DB 11; Length 479;
Best Local Similarity 62.5%; Pred. No. 5.5;
Matches 10; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 INWNGSGTGYADSVKG 17
| : : : : :
| : : : : :
| : : : : :
70 INSNGGTYSDTMKG 85

Db
RESULT 9
Q9UGP3 PRELIMINARY; PRT; 112 AA.
ID Q9UGP3
AC Q9UGP3;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Immunoglobulin heavy chain (fragment).
DE IGH.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Zafiroopoulos A., Kandilogianaki M., Dahlenborg C., Borraeack C.A.K.,

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Qy	1	GINWNGSTGYADSVK	17
	:	: ::	
Db	162	GTSYEGGNTGRPSVK	78
 RESULT 11			
Q7UUV9		PRELIMINARY:	PRT; 838 AA.
ID	Q7UUV9;		
AC	CACUUV9;		
DT	01-OCT-2003 (TrEMBLrel. 25; Created)		
DT	01-OCT-2003 (TrEMBLrel. 25; Last sequence update)		
DT	01-OCT-2003 (TrEMBLrel. 25; Last annotation update)		
DE	Hypothetical protein.		
DN	RB3034.		
OS	Rhodospirillum rubrum.		
OC	Bacteria; Planctomycetes; Planctomycetacia; Planctomycetales;		
OX	Planctomycetaceae; Pirellula.		
NCBI_TaxID=117;			
[1]_TaxID=117;			
RP	SEQUENCE FROM N.A.		
RC	STRAIN=1;		
RA	MEDLINE=22735913; PubMed=1285416;		
RA	Gloeckner F.O., Kube M., Bauer M., Teeling H., Lombardot T.,		
RA	Ludwig W., Gade D., Beck A., Borzym K., Heitmann K., Rabus R.,		
RA	Schlesner H., Amann R., Reinhardt R.;		
RT	"Complete genome sequence of the marine planctomycete Pirellula sp.		
RT	strain 1."		
RL	Proc. Natl. Acad. Sci. U.S.A. 100:8298-8303 (2003).		
KW	ENBL; BX294138; CAD72968.1; "		
DW	Hypothetical protein; Complete proteome.		
SEQ	SEQUENCE 838 AA; 94147 MW; 3321CFAA9E4A0336 CRC64;		
 Query Match 53.1%; Score 51; DB 16; Length 838; Best Local Similarity 60.0%; Pred No. 42; Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps			
Qy	2	GINWNGSTGYADSVK	16
	:	: ::	
Db	218	IKWQGTVAYATSVK	232
 RESULT 12			
ID	Q9U234	PRELIMINARY:	PRT; 767 AA.
ID	Q9U234;		
AC	CACUUV9;		
DT	01-MAY-2000 (TrEMBLrel. 13; Created)		
DT	01-MAY-2000 (TrEMBLrel. 13; Last sequence update)		
DT	01-MAY-2000 (TrEMBLrel. 13; Last sequence update)		
DE	Y56AJA.30 protein.		
DN	Y56AJA.30.		
OS	Caenorhabditis elegans.		
OC	Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditiida; Rhabditoidea;		
OC	Rhabditiidae; Peloderinae; Caenothabditis.		
NCBI_TaxID=6239;			
[1]			
RP	SEQUENCE FROM N.A.		
RA	Matthews L.;		
RA	Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.		
[2]			
RP	SEQUENCE FROM N.A.		
RC	MEDLINE=99069613; PubMed=9851916;		
RA	none;		
RT	"Genome sequence of the nematode C.elegans: A platform for		
RT	investigating biology.";		
RL	Science 282:2012-2018 (1998).		
DR	ENBL; AL132860; CAB60506.1; "		
DR	WormPep; Y56AJA.30; CE22595.		
SEQ	SEQUENCE 767 AA; 84076 MW; BB9D9E3C32E58CA08 CRC64;		
 Query Match 51.0%; Score 49; DB 5; Length 767; Best Local Similarity 47.1%; Pred. No. 77; Matches 8; Conservative 4; Mismatches 5; Indels 0; Gaps			

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QY 1 GINWGGSTGYADSVKG 17
   | : ||: ||| ||: |
Db 718 GYGGGTTGYMDNLVG 734

RESULT 13
Q9WZJ8 PRELIMINARY; PRT; 279 AA.
AC Q9WZJ8;
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein TM0739.
GN TM0739.
OS Thermotoga maritima
OC Bacteria; Thermotogae; Thermotogales; Thermotogaceae; Thermotoga.
OX NCBI_TaxID=2336;
RN [1]
RS SEQUENCE FROM N.A.
RC STRAIN=MSB8 / DSM 3109;
RC MEDLINE=99287316; PubMed=10360571;
RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
RA McDonald L., Utterback T.R., Malek J.A., Linher K.D., Garrett M.M.,
RA Stewart A.M., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,
RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
RT "Evidence for lateral gene transfer between Archaea and Bacteria from
RT genome sequence of the Thermotoga maritima.";
RL Nature 398:323-329(1999).
DR EMBL; AB001744; AAC35820.1; -.
DR PIR; F72339; F72339.
DR TIGR; TM0739; -.
DR GO; GO:0003824; F:catalytic activity; IEA.
DR InterPro; IPR006674; HD.
DR InterPro; IPR003607; Met_phosphohydro.
DR InterPro; IPR006675; Unchar_HDIG.
DR Pfam; PF01966; HD; 1.
DR SMART; SM00471; HDC; 1.
DR TIGRFAMs; TIGR00277; HDIG; 1.
DR Hypothetical protein; Complete proteome.
SQ SEQUENCE 279 AA; 32034 MW; BIC6494321C85B3B CRC64;

Query Match 50.0%; Score 48; DB 16; Length 279;
Best Local Similarity 60.0%; Pred. No. 34;
Matches 9; Conservative 2; Mismatches 2; Indels 2; Gaps 1;

QY 3 NWNGGSGTGYADSVKG 17
   | : ||: ||| ||: |
Db 200 NWDG--TCYPDGLKG 212

RESULT 14
Q9XGB4 PRELIMINARY; PRT; 298 AA.
AC Q9XGB4
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Chitinase.
OS Trifolium repens (Creeping white clover).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosid1; Fabales; Fabaceae; Papilionoideae; Trifolieae; Trifolium.
OX NCBI_TaxID=3899;
RN [1]
RS SEQUENCE FROM N.A.
RC STRAIN=cv. Dutch white; TISSUE=Root;
RA Crockard M.A., Broderick K., Pulvirenti G., Cooper J.E.;
RT "Expression of a class 3 chitinase in white clover roots following
RT homologous challenge.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.

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